

WEST Search History

DATE: Tuesday, October 15, 2002

<u>Set Name</u> side by side	<u>Query</u>	<u>Hit Count</u>	<u>Set Name</u> result set
<i>DB=USPT,PGPB,JPAB,EPAB,DWPI,TDBD; PLUR=YES; OP=OR</i>			
L17	L16 and (solid phase syntheses\$4) adj4 (peptide or polypeptide or dimer or cycl\$4)	18	L17
L16	L15 and (coupl\$4 or bridg\$4 or coupl\$4) adj4 (agent or moiety)	445	L16
L15	L14 and iminodiacetic adj4 acid	2227	L15
L14	s (di or tri or tetra) adj4 carboxylic adj4 acid	5188532	L14
L13	L12 and iminodiacetic adj4 acid	54	L13
L12	L11 and (coup\$6 or link\$6) adj6 (agent or technique)	608	L12
L11	L10 and ring adj5 structure	2074	L11
L10	L5 and dimer\$6	30286	L10
L9	L7 and achiral	1	L9
L8	L7 and imino adj4(di or tri or tetra)adj4 carboxylic adj4 acid	0	L8
L7	L6 and (aspartic or glutamic or glutaric)adj4 acid	525	L7
L6	L5 and iminodiacetic	1788	L6
L5	l1 and (solid phase synthesis)	868072	L5
L4	L2 and solid adj4 phase adj4 synthesis	26	L4
L3	L2 and achiral	8	L3
L2	L1 and (di or tri or tetra)adj4 carboxylic adj4 acid	9742	L2
L1	s borrelia	5179361	L1

END OF SEARCH HISTORY

FILE 'CA' ENTERED AT 11:01:02 ON 15 OCT 2002

L1 594 S (DI OR TRI OR TETRA) (5W) CARBOXY? (5W) ACID
L2 1 S L1 AND BORRELIA BURGDORFERI
L3 1 S L1 AND ACHIRAL
L4 0 S L1 NOT L2 AND L3
L5 594 S L1 NOT L2 OR L3
L6 593 S L1 NOT L2
L7 593 S L6 NOT L3

FILE 'REGISTRY' ENTERED AT 11:04:43 ON 15 OCT 2002

L8 7 S PVVAESPKKP/SQEP

FILE 'CA' ENTERED AT 11:05:45 ON 15 OCT 2002

L9 5 S L8
L10 1 S L1 AND BORRELIA

Run on: October 12, 2002, 20:43:46 ; Search time 30 Seconds

37.025 Million cell updates/sec

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Title: US-09-408-578A-1
Perfect score: 52
Sequence: 1 PVAESPCKP 10
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Gapop 10.0 , Gapext 0.5

```

Searched:      7475/4 segs, 111073796 residues
Total number of hits satisfying chosen parameters: 35207

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Minimum DB seq length: 0
Maximum DB seq length: 50
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Post-processing:	Minimum match	0%
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Listing first 45 summaries

Database :

1	/SIDS1/gcgdata/genseq/genseqp-emb1/AA1980.DAT.*
2	/SIDS1/gcgdata/genseq/genseqp-emb1/AA1981.DAT.*
3	/SIDS1/gcgdata/genseq/genseqp-emb1/AA1982.DAT.*
4	/SIDS1/gcgdata/genseq/genseqp-emb1/AA1983.DAT.*
5	/SIDS1/gcgdata/genseq/genseqp-emb1/AA1984.DAT.*
6	/SIDS1/gcgdata/genseq/genseqp-emb1/AA1985.DAT.*
7	/SIDS1/gcgdata/genseq/genseqp-emb1/AA1986.DAT.*
8	/SIDS1/gcgdata/genseq/genseqp-emb1/AA1987.DAT.*
9	/SIDS1/gcgdata/genseq/genseqp-emb1/AA1988.DAT.*
10	/SIDS1/gcgdata/genseq/genseqp-emb1/AA1989.DAT.*
11	/SIDS1/gcgdata/genseq/genseqp-emb1/AA1990.DAT.*
12	/SIDS1/gcgdata/genseq/genseqp-emb1/AA1991.DAT.*
13	/SIDS1/gcgdata/genseq/genseqp-emb1/AA1992.DAT.*
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16	/SIDS1/gcgdata/genseq/genseqp-emb1/AA1995.DAT.*
17	/SIDS1/gcgdata/genseq/genseqp-emb1/AA1996.DAT.*
18	/SIDS1/gcgdata/genseq/genseqp-emb1/AA1997.DAT.*
19	/SIDS1/gcgdata/genseq/genseqp-emb1/AA1998.DAT.*
20	/SIDS1/gcgdata/genseq/genseqp-emb1/AA1999.DAT.*
21	/SIDS1/gcgdata/genseq/genseqp-emb1/AA2000.DAT.*
22	/SIDS1/gcgdata/genseq/genseqp-emb1/AA2001.DAT.*

Pred. NO. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB	ID	Description
1	52	100.0	10	18	AAW41844	Modified B. burgdorferi
2	52	100.0	10	18	AAW41821	B. burgdorferi sensu stricto
3	52	100.0	11	18	AAW41825	Modified B. burgdorferi
4	52	100.0	15	18	AAW41827	B. burgdorferi sensu stricto
5	52	100.0	20	18	AAW41826	B. burgdorferi sensu stricto
6	52	100.0	23	20	AAV27428	B. burgdorferi sensu stricto
7	52	100.0	24	20	AAV27429	Borrelia outer surface protein
8	49	94.2	10	18	AAW41838	Borrelia outer surface protein
9	49	94.2	11	15	AAW70367	Modified B. burgdorferi
10	49	94.2	11	15	AAW70362	Borrelia ospC anti
11	48	92.3	10	18	AAW41841	Modified B. burgdorferi

[illegible]

ALIGNMENTS

XX	RESULT	1
XX	AAM41844	
ID	AAM41844	standard; peptide; 10 AA.
XX		
AC	AAM41844;	
DT	14-MAY-1998	(first entry)
DE	Modified B. burgdorferi sensu lato OspC C-terminal peptide	
XX	Sensu lato: outer surface protein C; OspC; diagnosis; Lyme	
KW	vaccine; infection.	
XX	Borrelia burgdorferi.	
OS	Synthetic.	
XX		
FH	Key	Location/Qualifiers
FT	Modified-site	10
FT	/label= amdated	
XX		
PN	W05742221-A1.	
PD	13-NOV-1997.	
XX		
PF	02-MAY-1997;	97WO-DK00203.
XX		
PR	02-MAY-1996;	96DK-0000526.
XX		
PA	(STAT-) SPATENS SERUMINSTITUT.	
PI	Holm A, Mathiesen MJ, Ostergaard S, Thølsen M,	
DR	WPI; 1997-558908/51.	

PT Detecting previous sensitisation to the OspC protein of Borrelia
 PT burgdorferi - by detecting immunoreactivity between patient T cells
 PT or immunoglobulins and C-terminal peptide of the protein

XX Example 3; Page 53; 95pp; English.

CC The present sequence was used in the development of a novel method
 CC for the identification of a patient's previous sensitisation to
 CC Borrelia burgdorferi sensu lato outer surface protein C (OspC).
 CC The method comprises reacting immunoglobulin (Ig) or T cells from
 CC the patient with a polypeptide of at most 60 amino acids containing
 CC a peptide with at least 50% identity to the B. burgdorferi derived
 CC sequence AAW41821, or its subsequences of at least 5 amino acids. The
 CC degree of immunological reactivity between the polypeptide and Ig
 CC or T cells is measured and significant reactivity is indicative of
 CC sensitisation.
 CC The method can be used to diagnose Lyme disease and is based on
 CC reactivity with antibodies against the OspC protein. The test can
 CC be done in vitro or in vivo, e.g. as a skin test. Vaccine
 CC compositions comprising the polypeptide can be used to protect
 CC humans and other animals against B. burgdorferi infection. The
 CC polypeptide provides higher sensitivity than full-length OspC, and
 CC so is better at detecting infection in its early stages, especially
 CC when combined with the known assay for flagellar proteins. The
 CC seven carboxy-terminal residues of AAW41821 represent an epitope
 CC essential for human immune response to OspC. The polypeptide is
 CC also easier to prepare and purify than (nearly) full-length
 CC protein, facilitating standardisation of the assay, and is less
 CC cross-reactive with antibodies raised against other antigens. The
 CC small size of the polypeptide allows a high density of binding
 CC sites to be created on a solid support. Incorporation of
 CC non-natural amino acid into the polypeptide increases its
 CC resistance to peptidases when used in vivo.

CC Sequence 10 AA:

Query Match 100.0%; Score 52; DB 18; Length 10;

Best Local Similarity 100.0%; Pred. No. 0.0078;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 PVVAESPCKP 10
 |||||
 Db 1 pvvaespkp 10

RESULT 2

AAW41821 standard; peptide; 10 AA.

AC AAW41821;

DT 14-MAY-1998 (first entry)

DE B. burgdorferi sensu lato OspC carboxy-terminal peptide.

KW Sensu lato; outer surface protein C; OspC; diagnosis; Lyme disease;
 KW vaccine; infection.

OS Borrelia burgdorferi.

PN WO9742221-A1.

PD 13-NOV-1997.

PF 02-MAY-1997; 97WO-DK00203.

PR 02-MAY-1996; 96DK-0000526.

PA (STAT-) STATENS SERUMINSTITUT.

PI Holm A, Mathiesen MJ, Ostergaard S, Thiesen M;

DR WPI: 1997-558908/51.

XX Detecting previous sensitisation to the OspC protein of Borrelia
 PT burgdorferi - by detecting immunoreactivity between patient T cells
 PT or immunoglobulins and C-terminal peptide of the protein

XX Claim 1; Page 77; 95pp; English.

CC The present sequence was used in the development of a novel method
 CC for the identification of a patient's previous sensitisation to
 CC Borrelia burgdorferi sensu lato outer surface protein C (OspC).
 CC The method comprises reacting immunoglobulin (Ig) or T cells from
 CC the patient with a polypeptide of at most 60 amino acids containing
 CC a peptide with at least 50% identity to the B. burgdorferi derived
 CC sequence AAW41821, or its subsequences of at least 5 amino acids. The
 CC degree of immunological reactivity between the polypeptide and Ig
 CC or T cells is measured and significant reactivity is indicative of
 CC sensitisation.
 CC The method can be used to diagnose Lyme disease and is based on
 CC reactivity with antibodies against the OspC protein. The test can
 CC be done in vitro or in vivo, e.g. as a skin test. Vaccine
 CC compositions comprising the polypeptide can be used to protect
 CC humans and other animals against B. burgdorferi infection. The
 CC polypeptide provides higher sensitivity than full-length OspC, and
 CC so is better at detecting infection in its early stages, especially
 CC when combined with the known assay for flagellar proteins. The
 CC seven carboxy-terminal residues of AAW41821 represent an epitope
 CC essential for human immune response to OspC. The polypeptide is
 CC also easier to prepare and purify than (nearly) full-length
 CC protein, facilitating standardisation of the assay, and is less
 CC cross-reactive with antibodies raised against other antigens. The
 CC small size of the polypeptide allows a high density of binding
 CC sites to be created on a solid support. Incorporation of
 CC non-natural amino acid into the polypeptide increases its
 CC resistance to peptidases when used in vivo.

CC Sequence 10 AA:

Query Match 100.0%; Score 52; DB 18; Length 10;

Best Local Similarity 100.0%; Pred. No. 0.0078;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 PVVAESPCKP 10
 |||||
 Db 1 pvvaespkp 10

RESULT 3

AAW41825 standard; peptide; 11 AA.

AC AAW41825;

DT 14-MAY-1998 (first entry)

DE Modified B. burgdorferi sensu lato OspC C-terminal peptide.

KW Sensu lato; outer surface protein C; OspC; diagnosis; Lyme disease;
 KW vaccine; infection.

OS Borrelia burgdorferi.

PN WO9742221-A1.

PD 13-NOV-1997.

PF 02-MAY-1997; 97WO-DK00203.

PA (STAT-) STATENS SERUMINSTITUT.

PI Holm A, Mathiesen MJ, Ostergaard S, Thiesen M;

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XX 02-MAY-1996; 96DK-0000526.
XX (STAT-) STATENS SERUMINSTITUT.
XX Holm A, Mathiesen MJ, Ostergaard S, Theisen M;
XX WPI; 1997-558908/51.
XX
XX Detecting previous sensitisation to the OspC protein of Borrelia
XX burgdorferi - by detecting immunoreactivity between patient T cells
XX or immunoglobulins and C-terminal peptide of the protein
XX
XX Example; Page 40; 95pp; English.
XX
XX The present sequence was used in the development of a novel method
XX for the identification of a patient's previous sensitisation to
XX Borrelia burgdorferi sensu lato outer surface protein C (OspC).
XX The method comprises reacting immunoglobulin (Ig) or T cells from
XX the patient with a polypeptide of at most 60 amino acids containing
XX a peptide with at least 50% identity to the B. burgdorferi derived
XX sequence AAW41821, or its subsequences of at least 5 amino acids. The
XX degree of immunological reactivity between the polypeptide and Ig
XX or T cells is measured and significant reactivity is indicative of
XX sensitisation.
XX The method can be used to diagnose Lyme disease and is based on
XX reactivity with antibodies against the OspC protein. The test can
XX be done in vitro or in vivo, e.g. as a skin test. Vaccine
XX compositions comprising the polypeptide can be used to protect
XX humans and other animals against B. burgdorferi infection. The
XX polypeptide provides higher sensitivity than full-length OspC, and
XX so is better at detecting infection in its early stages, especially
XX when combined with the known assay for flagellar proteins. The
XX seven carboxy-terminal residues of AAW41821 represent an epitope
XX essential for human immune response to OspC. The polypeptide is
XX also easier to prepare and purify than (nearly) full-length
XX protein, facilitating standardisation of the assay, and is less
XX cross-reactive with antibodies raised against other antigens. The
XX small size of the polypeptide allows a high density of binding
XX sites to be created on a solid support. Incorporation of
XX non-natural amino acid into the polypeptide increases its
XX resistance to peptidases when used in vivo.
XX
XX Sequence 11 AA:
SO

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Query Match 100.0%; Score 52; DB 18; Length 11;
Best Local Similarity 100.0%; Pred. No. 0.0085;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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OY 1 PVVAESPKRP 10
    |||||
    2 PVVAESPKRP 11

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RESULT 4
AAW41827 standard; peptide; 15 AA.
XX AAW41827;
XX AC
XX 14-MAY-1998 (first entry)
XX DT
XX B. burgdorferi sensu lato OspC carboxy-terminal peptide.
XX DE
XX Sensu lato; outer surface protein C; OspC; diagnosis; Lyme disease;
XX KW vaccine; infection.
XX OS Borrelia burgdorferi.
XX PN W09742221-A1.
XX PD 13-NOV-1997.

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XX 02-MAY-1997; 97WO-DK00203.
XX (STAT-) STATENS SERUMINSTITUT.
XX Holm A, Mathiesen MJ, Ostergaard S, Theisen M;
XX WPI; 1997-558908/51.
XX
XX Detecting previous sensitisation to the OspC protein of Borrelia
XX burgdorferi - by detecting immunoreactivity between patient T cells
XX or immunoglobulins and C-terminal peptide of the protein
XX
XX Example 3; Page 51; 95pp; English.
XX
XX The present sequence was used in the development of a novel method
XX for the identification of a patient's previous sensitisation to
XX Borrelia burgdorferi sensu lato outer surface protein C (OspC).
XX The method comprises reacting immunoglobulin (Ig) or T cells from
XX the patient with a polypeptide of at most 60 amino acids containing
XX a peptide with at least 50% identity to the B. burgdorferi derived
XX sequence AAW41821, or its subsequences of at least 5 amino acids. The
XX degree of immunological reactivity between the polypeptide and Ig
XX or T cells is measured and significant reactivity is indicative of
XX sensitisation.
XX The method can be used to diagnose Lyme disease and is based on
XX reactivity with antibodies against the OspC protein. The test can
XX be done in vitro or in vivo, e.g. as a skin test. Vaccine
XX compositions comprising the polypeptide can be used to protect
XX humans and other animals against B. burgdorferi infection. The
XX polypeptide provides higher sensitivity than full-length OspC, and
XX so is better at detecting infection in its early stages, especially
XX when combined with the known assay for flagellar proteins. The
XX seven carboxy-terminal residues of AAW41821 represent an epitope
XX essential for human immune response to OspC. The polypeptide is
XX also easier to prepare and purify than (nearly) full-length
XX protein, facilitating standardisation of the assay, and is less
XX cross-reactive with antibodies raised against other antigens. The
XX small size of the polypeptide allows a high density of binding
XX sites to be created on a solid support. Incorporation of
XX non-natural amino acid into the polypeptide increases its
XX resistance to peptidases when used in vivo.
XX
XX Sequence 15 AA:
SO

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Query Match 100.0%; Score 52; DB 18; Length 15;
Best Local Similarity 100.0%; Pred. No. 0.012;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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OY 1 PVVAESPKRP 10
    |||||
    6 PVVAESPKRP 15

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RESULT 5
AAW41826 standard; peptide; 20 AA.
XX AAW41826;
XX AC
XX 14-MAY-1998 (first entry)
XX DT
XX B. burgdorferi sensu lato OspC carboxy-terminal peptide.
XX DE
XX Sensu lato; outer surface protein C; OspC; diagnosis; Lyme disease;
XX KW vaccine; infection.
XX OS Borrelia burgdorferi.
XX PN W09742221-A1.

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XX 13-NOV-1997.
 PD
 XX
 PF 02-MAY-1997; 97WO-DK00203.
 PR 02-MAY-1996; 96DK-0000526.
 XX
 PA (STAT-) STATENS SERUMINSTITUT.
 XX
 PI Holm A, Mathiesen MJ, Ostergaard S, Thelsen M;
 DR WPI; 1997-558908/51.
 XX
 PT Detecting previous sensitisation to the OspC protein of *Borrelia*
 XX burgdorferi - by detecting immunoreactivity between patient T cells
 XX or immunoglobulins and C-terminal peptide of the protein
 PS
 PS Example 3; Page 51; 95pp; English.
 CC
 CC The present sequence was used in the development of a novel method
 CC for the identification of a patient's previous sensitisation to
 CC *Borrelia burgdorferi* sensu lato outer surface protein C (OspC).
 CC The method comprises reacting immunoglobulin (Ig) or T cells from
 CC the patient with a polypeptide of at most 60 amino acids containing
 CC a peptide with at least 50% identity to the B. burgdorferi derived
 CC sequence AAW41821, or its subsequences of at least 5 amino acids. The
 CC degree of immunological reactivity between the polypeptide and Ig
 CC or T cells is measured and significant reactivity is indicative of
 CC sensitisation.
 CC The method can be used to diagnose Lyme disease and is based on
 CC reactivity with antibodies against the OspC protein. The test can
 CC be done in vitro or in vivo, e.g. as a skin test. Vaccine
 CC compositions comprising the polypeptide can be used to protect
 CC humans and other animals against B. burgdorferi infection. The
 CC polypeptide provides higher sensitivity than full-length OspC, and
 CC so is better at detecting infection in its early stages, especially
 CC when combined with the known assay for flagellar proteins. The
 CC seven carboxy-terminal residues of AAW41821 represent an epitope
 CC essential for human immune response to OspC. The polypeptide is
 CC also easier to prepare and purify than (nearly) full-length
 CC protein, facilitating standardisation of the assay, and is less
 CC cross-reactive with antibodies raised against other antigens. The
 CC small size of the polypeptide allows a high density of binding
 CC sites to be created on a solid support. Incorporation of
 CC non-natural amino acid into the polypeptide increases its
 CC resistance to peptidases when used in vivo.
 XX
 SQ Sequence 20 AA;
 XX
 XX
 Query Match 100.0%; Score 52; DB 18; Length 20;
 Best Local Similarity 100.0%; PId. NO. 0.016;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0
 QY 1 PVAESPCKP 10
 | | | | | | | | | |
 Db 11 pvaespckp 20
 XX
 RESULT 6
 AAY27428
 ID AAY27428 standard; peptide: 23 AA.
 XX
 AC AAY27428;
 XX
 DT 26-NOV-1999 (first entry)
 XX
 DE *Borrelia* outer surface protein C (OspC) C-terminal peptide fragment.
 XX
 KW *Borrelia*: Igm antibody; outer surface protein C; OspC; deer tick;
 KW cerebrospinal fluid; Lyme borreliosis; micro-capture assay; flagellum;
 XX epitope.
 XX

Key	Location/Qualifiers
Modified-site	1 /note= "linked to biotin via an O-linker of formula [2-aminoethoxy]ethoxy acetic acid"
EP949508-A1	
13-OCT-1999.	
07-APR-1999;	99EP-0610026.
08-APR-1998;	98DK-0000516.
(DAKO-) DAKO AS.	
Staffeldt Schou O, Winther L, Stender H;	
MP1: 1999-553537/47	
Diagnosing Lyme-borreliosis by detecting antibodies against two antigens simultaneously -	
Example 1; Page 7; 23pp; English.	
The invention provides a new method for detecting Igm antibodies against Borrelia burgdorferi in a sample of human or animal fluid. The method comprises: (1) contacting antibodies in the sample with anti-igm immobilized to a solid support, (2) separating the support from the liquid phase; and (3) contacting the bound antibodies with a complex comprising at least one set of B. burgdorferi (outer surface protein C (OspC) peptides and/or at least one set of other B. burgdorferi peptides, each attached to a carrier; and (4) detecting the presence of antibodies against B. burgdorferi. The new method may be used to detect antibodies against B. burgdorferi in (especially) serum or cerebrospinal fluid samples from patients bitten by deer ticks. B. burgdorferi causes Lyme borreliosis so detection of antibodies against it allows diagnosis of infection by this organism. The method is a micro-capture assay in which the antigen complex is a combination of the B. burgdorferi flagellum and OspC peptides. The presence of epitopes from both antigens in the complex allows the simultaneous detection of serum antibodies against these proteins which increases the sensitivity of the test. The two antigens are pure, which also decreases the possibility of cross-reactivity. The present sequence represents a Borrelia OspC C-terminal peptide fragment.	
Sequence	23 AA;
Query Match	100.0%; Score 52; DB 20; Length 23;
Best Local Similarity	100.0%; Pred. No. 0.018;
Matches	10; Conservative 0; Mismatches 0; Indels 0; Gaps 0
QY	1 PVVAESP KKP 10
DB	14 PVVAESP KKP 23
RESULT	7
AAV27429	
AAV27429	standard; peptide; 24 AA.
AAV27429;	
26-NOV-1999	(first entry)
Borrelia outer surface protein C (OspC) C-terminal peptide fragment.	
Borrelia; Igm antibody; outer surface protein C; OspC; deer tick; cerebrospinal fluid; Lyme borreliosis; micro-capture assay; flagellum; epitope.	

OS Synthetic.
 OS Borrelia burgdorferi.
 PN EP949508-A1.
 XX
 PD 13-OCT-1999.
 XX
 PF 07-APR-1999; 99EP-0610026.
 XX
 PR 08-APR-1998; 98DK-0000516.
 XX
 PA (DAKO-) DAKO AS.
 PI Staffeldt Schou O, Winther L, Stender H;
 DR WPI; 1999-553537/47.
 XX
 PT Diagnosing Lyme borreliosis by detecting antibodies against two
 PT antigens simultaneously -
 XX
 PS Example 1; Page 7; 23pp; English.
 CC The invention provides a new method for detecting IGM antibodies against
 CC Borrelia burgdorferi in a sample of human or animal fluid. The method
 CC comprises: (1) contacting antibodies in the sample with anti-IGM
 CC immobilized to a solid support, (2) separating the support from the
 CC liquid phase; and (3) contacting the bound antibodies with a complex
 CC comprising at least one set of B. burgdorferi (outer surface protein C
 CC (OspC) peptides and/or at least one set of other B. burgdorferi peptides,
 CC each attached to a carrier; and (4) detecting the presence of antibodies
 CC against B. burgdorferi. The new method may be used to detect antibodies
 CC against B. burgdorferi in (especially) serum or cerebrospinal fluid
 CC samples from patients bitten by deer ticks. B. burgdorferi causes Lyme
 CC borreliosis so detection of antibodies against it allows diagnosis of
 CC infection by this organism. The method is a micro-capture assay in which
 CC the antigen complex is a combination of the B. burgdorferi flagellum and
 CC OspC peptides. The presence of epitopes from both antigens in the complex
 CC allows the simultaneous detection of serum antibodies against these
 CC proteins which increases the sensitivity of the test. The two antigens
 CC are pure, which also decreases the possibility of cross-reactivity. The
 CC present sequence represents a Borrelia OspC C-terminal peptide fragment,
 CC where the N-terminal cysteine residue has been incorporated to provide a
 CC SH group to be used in a coupling reaction.
 CC
 CC Sequence 24 AA:
 SQ
 Query Match 100.0%; Score 52; DB 20; Length 24;
 Best Local Similarity 100.0%; Pred. No. 0.019; 0; Indels 0; Gaps 0;
 Matches 10; Conservative 0; Mismatches 0;
 OY 1 PVVAESPCKP 10
 |||||
 DB 15 pvaespckp 24
 RESULT 8
 AAM41838 standard; peptide; 10 AA.
 AC AAM41838;
 XX
 DT 14-MAY-1998 (first entry)
 XX
 DE Modified B. burgdorferi sensu lato OspC C-terminal peptide.
 XX
 DE Senu lato; outer surface protein C; OspC; diagnosis; Lyme disease;
 KW vaccine; infection.
 XX
 XX Borrelia burgdorferi.
 OS Synthetic.
 OS
 PN MO9742221-A1.

XX
 PD 13-NOV-1997.
 XX
 XX 02-MAY-1997; 97WO-DK00203.
 PF
 XX
 PR 02-MAY-1996; 96DK-0000526.
 XX
 PA (STAT-) STATENS SERUMINSTITUT.
 XX
 PI Holm A, Mathiesen MJ, Ostergaard S, Theisen M;
 DR WPI; 1997-558908/51.
 XX
 PT Detecting previous sensitisation to the OspC protein of Borrelia
 PT burgdorferi - by detecting immunoreactivity between patient T cells
 PT or immunoglobulins and C-terminal peptide of the protein
 XX
 PS Example 3; Page 52; 95pp; English.
 CC The present sequence was used in the development of a novel method
 CC for the identification of a patient's previous sensitisation to
 CC Borrelia burgdorferi sensu lato outer surface protein C (OspC).
 CC The method comprises reacting immunoglobulin (Ig) or T cells from
 CC the patient with a polypeptide of at most 60 amino acids containing
 CC a peptide with at least 50% identity to the B. burgdorferi derived
 CC sequence AAM41821, or its subsequences of at least 5 amino acids. The
 CC degree of immunological reactivity between the polypeptide and Ig
 CC or T cells is measured and significant reactivity is indicative of
 CC sensitisation.
 CC The method can be used to diagnose Lyme disease and is based on
 CC reactivity with antibodies against the OspC protein. The test can
 CC be done in vitro or in vivo, e.g. as a skin test. Vaccine
 CC compositions comprising the polypeptide can be used to protect
 CC humans and other animals against B. burgdorferi infection. The
 CC polypeptide provides higher sensitivity than full-length OspC, and
 CC so is better at detecting infection in its early stages, especially
 CC when combined with the known assay for flagellar proteins. The
 CC seven carboxy-terminal residues of AAM41821 represent an epitope
 CC essential for human immune response to OspC. The polypeptide is
 CC also easier to prepare and purify than (nearly) full-length
 CC protein, facilitating standardisation of the assay, and is less
 CC cross-reactive with antibodies raised against other antigens. The
 CC small size of the polypeptide allows a high density of binding
 CC sites to be created on a solid support. Incorporation of
 CC non-natural amino acid into the polypeptide increases its
 CC resistance to peptidases when used in vivo.
 CC
 CC Sequence 10 AA:
 SQ
 Query Match 94.2%; Score 49; DB 18; Length 10;
 Best Local Similarity 90.0%; Pred. No. 0.026;
 Matches 9; Conservative 1; Mismatches 0; Indels 0; Gaps 0;
 OY 1 PVVAESPCKP 10
 |||||
 DB 1 pvaespckp 10
 RESULT 9
 AAR70367
 ID AAR70367 standard; peptide; 11 AA.
 AC AAR70367;
 XX
 DT 25-MAY-1995 (first entry)
 XX
 DE Borrelia OspC antigen epitope.
 XX
 DE OspC antigen; vaccine; Lyme disease; borreliosis; immunogen;
 KW serovar typing; restriction fragment length polymorphism;
 KW RFLP.
 XX

OS Borrelia burgdorferi.
 XX
 PN WO9425596-A.
 XX
 PD 10-NOV-1994.
 XX
 PF 29-APR-1994; 94MO-EP01365.
 XX
 PR 29-APR-1993; 93US-0053863.
 XX
 PA (IMMO) IMMUNO AG.
 XX
 PI Crowe B, Dornier F, Lavey I;
 XX
 DR WPI: 1994-358273/44.
 XX
 PT Immunogenic composition comprising OspC antigens - for the
 PT treatment of Lyme borreliosis in different, specific geographical
 PT areas.
 PS Claim 19; Page 57; 115pp; English.
 CC
 CC A vaccine for Lyme disease includes selected OspC antigen
 CC formulations based on defined OspC families resolved by serovar
 CC typing and RFLP typing of strains of worldwide origin. The
 CC antigens comprise 1 or more of the epitopes given in AAR70360-69
 CC or their variants or mimetics.
 XX
 SQ Sequence 11 AA;
 OY
 Query Match 94.2%; Score 49; DB 15; Length 11;
 Best Local Similarity 90.0%; Pred. No. 0.029;
 Matches 9; Conservative 1; Mismatches 0; Indels 0; Gaps 0;
 DB 1 PVVASEPKKP 10
 2 PVVASEPKKP 11
 RESULT 10
 AAR70362
 ID AAR70362 standard; peptide; 15 AA.
 AC AAR70362;
 XX
 DT 25-MAY-1995 (first entry)
 XX
 DE Borrelia OspC antigen epitope.
 XX
 KW OspC antigen; vaccine; Lyme disease; borreliosis; immunogen;
 KW serovar typing; restriction fragment length polymorphism;
 KW RFLP.
 XX
 OS Borrelia burgdorferi.
 XX
 PN WO9425596-A.
 XX
 PD 10-NOV-1994.
 XX
 PF 29-APR-1994; 94MO-EP01365.
 XX
 PR 29-APR-1993; 93US-0053863.
 XX
 PA (IMMO) IMMUNO AG.
 XX
 PI Crowe B, Dornier F, Lavey I;
 XX
 DR WPI: 1994-358273/44.
 XX
 PT Immunogenic composition comprising OspC antigens - for the
 PT treatment of Lyme borreliosis in different, specific geographical
 PT areas.

XX
 PS Claim 19; Page 56; 115pp; English.
 XX
 CC A vaccine for Lyme disease includes selected OspC antigen
 CC formulations based on defined OspC families resolved by serovar
 CC typing and RFLP typing of strains of worldwide origin. The
 CC antigens comprise 1 or more of the epitopes given in AAR70360-69
 CC or their variants or mimetics.
 XX
 SQ Sequence 15 AA;
 OY
 Query Match 94.2%; Score 49; DB 15; Length 15;
 Best Local Similarity 90.0%; Pred. No. 0.038;
 Matches 9; Conservative 1; Mismatches 0; Indels 0; Gaps 0;
 DB 1 PVVASEPKKP 10
 6 PVVASEPKKP 15
 RESULT 11
 AAW41841
 ID AAW41841 standard; peptide; 10 AA.
 AC AAW41841;
 XX
 DT 14-MAY-1998 (first entry)
 XX
 DE Modified B. burgdorferi sensu lato OspC C-terminal peptide.
 XX
 KW Senu lato; outer surface protein C; OspC; diagnosis; Lyme disease;
 KW vaccine; infection.
 XX
 OS Borrelia burgdorferi.
 XX
 PN WO9742221-A1.
 XX
 PD 13-NOV-1997.
 XX
 PF 02-MAY-1997; 97WO-DK00203.
 XX
 PR 02-MAY-1996; 96DK-0000526.
 XX
 PA (STAT-) STATENS SERUMINSTITUT.
 XX
 PI Holm A, Mathiesen MJ, Ostergaard S, Theisen M;
 XX
 DR WPI: 1997-558908/51.
 XX
 PT Detecting previous sensitisation to the OspC protein of Borrelia
 PT burgdorferi - by detecting immunoreactivity between patient T cells
 PT or immunoglobulins and C-terminal peptide of the protein
 XX
 PS Example 3; Page 52; 95pp; English.
 CC
 CC The present sequence was used in the development of a novel method
 CC for the identification of a patient's previous sensitisation to
 CC Borrelia burgdorferi sensu lato outer surface protein C (OspC).
 CC The method comprises reacting immunoglobulin (Ig) or T cells from
 CC the patient with a polypeptide of at most 60 amino acids containing
 CC a peptide with at least 50% identity to the B. burgdorferi derived
 CC sequence AAW41821, or its subsequences of at least 5 amino acids. The
 CC degree of immunological reactivity between the polypeptide and Ig
 CC or T cells is measured and significant reactivity is indicative of
 CC sensitisation.
 CC The method can be used to diagnose Lyme disease and is based on
 CC reactivity with antibodies against the OspC protein. The test can
 CC be done in vitro or in vivo, e.g. as a skin test. Vaccine
 CC compositions comprising the polypeptide can be used to protect
 CC humans and other animals against B. burgdorferi infection. The
 CC polypeptide provides higher sensitivity than full-length OspC, and

so is better at detecting infection in its early stages, especially when combined with the known assay for flagellar proteins. The seven carboxy-terminal residues of AAW41821 represent an epitope essential for human immune response to OspC. The polypeptide is also easier to prepare and purify than (nearly) full-length protein, facilitating standardisation of the assay, and is less cross-reactive with antibodies raised against other antigens. The small size of the polypeptide allows a high density of binding sites to be created on a solid support. Incorporation of non-natural amino acid into the polypeptide increases its resistance to peptidases when used in vivo.

Sequence 10 AA:

Query Match 92.3%; Score 48; DB 18; Length 10;
Best Local Similarity 90.0%; Pred. No. 0.039;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 1 PVVAESPCKP 10
1111111111
DB 1 pvaeespkxp 10

RESULT 12
AAW41842
ID AAW41842 standard; peptide; 10 AA.

AAW41842;

14-MAY-1998 (first entry)

Modified B. burgdorferi sensu lato OspC C-terminal peptide.

Sensu lato; outer surface protein C; OspC; diagnosis; Lyme disease;

vaccine; Infection.

Borrelia burgdorferi.

Synthetic.

MO9742221-A1.

13-NOV-1997.

02-MAY-1997; 97WO-DK00203.

02-MAY-1996; 96DK-0000526.

(STAT-) STATENS SERUMINSTITUT.

Holm A, Mathiesen MJ, Ostergaard S, Theisen M;

WPI; 1997-558908/51.

Detecting previous sensitisation to the OspC protein of Borrelia burgdorferi - by detecting immunoreactivity between patient T cells or immunoglobulins and C-terminal peptide of the protein

Example 3; Page 52; 95pp; English.

The present sequence was used in the development of a novel method for the identification of a patient's previous sensitisation to Borrelia burgdorferi sensu lato outer surface protein C (OspC). The method comprises reacting immunoglobulin (Ig) or T cells from the patient with a polypeptide of at most 60 amino acids containing a peptide with at least 50% identity to the B. burgdorferi derived sequence AAW41821, or its subsequences of at least 5 amino acids. The degree of immunological reactivity between the polypeptide and Ig or T cells is measured and significant reactivity is indicative of sensitisation.

The method can be used to diagnose Lyme disease and is based on reactivity with antibodies against the OspC protein. The test can be done in vitro or in vivo, e.g. as a skin test. Vaccine

compositions comprising the polypeptide can be used to protect humans and other animals against B. burgdorferi infection. The polypeptide provides higher sensitivity than full-length OspC, and so is better at detecting infection in its early stages, especially when combined with the known assay for flagellar proteins. The seven carboxy-terminal residues of AAW41821 represent an epitope essential for human immune response to OspC. The polypeptide is also easier to prepare and purify than (nearly) full-length protein, facilitating standardisation of the assay, and is less cross-reactive with antibodies raised against other antigens. The small size of the polypeptide allows a high density of binding sites to be created on a solid support. Incorporation of non-natural amino acid into the polypeptide increases its resistance to peptidases when used in vivo.

Sequence 10 AA:

Query Match 92.3%; Score 48; DB 18; Length 10;
Best Local Similarity 90.0%; Pred. No. 0.039;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 1 PVVAESPCKP 10
1111111111
DB 1 pvaeespkxp 10

RESULT 13
AAW41848
ID AAW41848 standard; peptide; 10 AA.

AAW41848;

14-MAY-1998 (first entry)

Modified B. burgdorferi sensu lato OspC C-terminal peptide.

Sensu lato; outer surface protein C; OspC; diagnosis; Lyme disease;

vaccine; Infection.

Borrelia burgdorferi.

Synthetic.

MO9742221-A1.

13-NOV-1997.

02-MAY-1997; 97WO-DK00203.

02-MAY-1996; 96DK-0000526.

(STAT-) STATENS SERUMINSTITUT.

Holm A, Mathiesen MJ, Ostergaard S, Theisen M;

WPI; 1997-558908/51.

Detecting previous sensitisation to the OspC protein of Borrelia burgdorferi - by detecting immunoreactivity between patient T cells or immunoglobulins and C-terminal peptide of the protein

Example 3; Page 54; 95pp; English.

The present sequence was used in the development of a novel method for the identification of a patient's previous sensitisation to Borrelia burgdorferi sensu lato outer surface protein C (OspC). The method comprises reacting immunoglobulin (Ig) or T cells from the patient with a polypeptide of at most 60 amino acids containing a peptide with at least 50% identity to the B. burgdorferi derived sequence AAW41821, or its subsequences of at least 5 amino acids. The degree of immunological reactivity between the polypeptide and Ig or T cells is measured and significant reactivity is indicative of sensitisation.

CC The method can be used to diagnose Lyme disease and is based on
 CC reactivity with antibodies against the OspC protein. The test can
 CC be done in vitro or in vivo, e.g. as a skin test. Vaccine
 CC compositions comprising the polypeptide can be used to protect
 CC humans and other animals against B. burgdorferi infection. The
 CC polypeptide provides higher sensitivity than full-length OspC, and
 CC so is better at detecting infection in its early stages, especially
 CC when combined with the known assay for flagellar proteins. The
 CC seven carboxy-terminal residues of AAM41821 represent an epitope
 CC essential for human immune response to OspC. The polypeptide is
 CC also easier to prepare and purify than (nearly) full-length
 CC protein, facilitating standardisation of the assay, and is less
 CC cross-reactive with antibodies raised against other antigens. The
 CC small size of the polypeptide allows a high density of binding
 CC sites to be created on a solid support. Incorporation of
 CC non-natural amino acid into the polypeptide increases its
 CC resistance to peptidases when used in vivo.

CC Sequence 10 AA;

Query Match 90.4%; Score 47; DB 18; Length 10;
 Best Local Similarity 90.0%; Pred. No. 0.059;
 Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 PVVAESPCKP 10
 ||| |||||
 DB 1 pvvspckp 10

RESULT 14

AAM41849
 ID AAM41849 standard; peptide; 10 AA.

XX AAM41849;

DT 14-MAY-1998 (first entry)

DE Modified B. burgdorferi sensu lato OspC C-terminal peptide.

KW Sensu lato; outer surface protein C; OspC; diagnosis; Lyme disease;

KM vaccine; infection.

OS Borrelia burgdorferi.

OS Synthetic.

PN WO9742221-A1.

PD 13-NOV-1997.

PF 02-MAY-1997; 97WO-DK00203.

PR 02-MAY-1996; 96DK-0000526.

PA (STAT-) STATENS SERUMINSTITUTT.

PI Holm A, Mathiesen MJ, Ostergaard S, Theisen M;

DR WPI; 1997-558908/51.

PT Detecting previous sensitisation to the OspC protein of Borrelia
 PT burgdorferi - by detecting immunoreactivity between patient T cells
 PT or immunoglobulins and C-terminal peptide of the protein

XX Example 3; Page 54; 95pp; English.

CC The present sequence was used in the development of a novel method
 CC for the identification of a patient's previous sensitisation to
 CC Borrelia burgdorferi sensu lato outer surface protein C (OspC).
 CC The method comprises reacting immunoglobulin (Ig) or T cells from
 CC the patient with a polypeptide of at most 60 amino acids containing
 CC a peptide with at least 50% identity to the B. burgdorferi derived
 CC sequence AAM41821, or its subsequences of at least 5 amino acids. The

CC degree of immunological reactivity between the polypeptide and Ig
 CC or T cells is measured and significant reactivity is indicative of
 CC sensitisation.
 CC The method can be used to diagnose Lyme disease and is based on
 CC reactivity with antibodies against the OspC protein. The test can
 CC be done in vitro or in vivo, e.g. as a skin test. Vaccine
 CC compositions comprising the polypeptide can be used to protect
 CC humans and other animals against B. burgdorferi infection. The
 CC polypeptide provides higher sensitivity than full-length OspC, and
 CC so is better at detecting infection in its early stages, especially
 CC when combined with the known assay for flagellar proteins. The
 CC seven carboxy-terminal residues of AAM41821 represent an epitope
 CC essential for human immune response to OspC. The polypeptide is
 CC also easier to prepare and purify than (nearly) full-length
 CC protein, facilitating standardisation of the assay, and is less
 CC cross-reactive with antibodies raised against other antigens. The
 CC small size of the polypeptide allows a high density of binding
 CC sites to be created on a solid support. Incorporation of
 CC non-natural amino acid into the polypeptide increases its
 CC resistance to peptidases when used in vivo.

CC Sequence 10 AA;

Query Match 90.4%; Score 47; DB 18; Length 10;
 Best Local Similarity 90.0%; Pred. No. 0.059;
 Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 PVVAESPCKP 10
 ||| |||||
 DB 1 pvvspckp 10

RESULT 15

AAM41845
 ID AAM41845 standard; peptide; 10 AA.

XX AAM41845;

DT 14-MAY-1998 (first entry)

DE Modified B. burgdorferi sensu lato OspC C-terminal peptide.

KW Sensu lato; outer surface protein C; OspC; diagnosis; Lyme disease;

KM vaccine; infection.

OS Borrelia burgdorferi.

OS Synthetic.

PN WO9742221-A1.

PD 13-NOV-1997.

PF 02-MAY-1997; 97WO-DK00203.

PR 02-MAY-1996; 96DK-0000526.

PA (STAT-) STATENS SERUMINSTITUTT.

PI Holm A, Mathiesen MJ, Ostergaard S, Theisen M;

DR WPI; 1997-558908/51.

PT Detecting previous sensitisation to the OspC protein of Borrelia
 PT burgdorferi - by detecting immunoreactivity between patient T cells
 PT or immunoglobulins and C-terminal peptide of the protein

XX Example 3; Page 54; 95pp; English.

PS Example 3: Page 53; 95pp; English.

CC The present sequence was used in the development of a novel method
CC for the identification of a patient's previous sensitisation to
CC *Borellia burgdorferi* sensu lato outer surface protein C (OspC).
CC The method comprises reacting immunoglobulin (Ig) or T cells from
CC the patient with a polypeptide of at most 60 amino acids containing
CC a peptide with at least 50% identity to the *B. burgdorferi* derived
CC sequence AAW41821, or its subsequences of at least 5 amino acids. The
CC degree of immunological reactivity between the polypeptide and Ig
CC or T cells is measured and significant reactivity is indicative of
CC sensitisation.
CC The method can be used to diagnose Lyme disease and is based on
CC reactivity with antibodies against the OspC protein. The test can
CC be done in vitro or in vivo, e.g. as a skin test. Vaccine
CC compositions comprising the polypeptide can be used to protect
CC humans and other animals against *B. burgdorferi* infection. The
CC polypeptide provides higher sensitivity than full-length OspC, and
CC so is better at detecting infection in its early stages, especially
CC when combined with the known assay for flagellar proteins. The
CC seven carboxy-terminal residues of AAW41821 represent an epitope
CC essential for human immune response to OspC. The polypeptide is
CC also easier to prepare and purify than (nearly) full-length
CC protein, facilitating standardisation of the assay, and is less
CC cross-reactive with antibodies raised against other antigens. The
CC small size of the polypeptide allows a high density of binding
CC sites to be created on a solid support. Incorporation of
CC non-natural amino acid into the polypeptide increases its
CC resistance to peptidases when used in vivo.

XX
SQ Sequence 10 AA;

Query Match

Best Local Similarity 88.5%; Score 46; DB 18; Length 10;

Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

OY 1 PVVAESPCKP 10
|||
Db 1 pvvaespkxp 10

Search completed: October 12, 2002, 20:48:43
Job time: 297 sec

GenCore version 4.5
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OM protein - protein search, using sw model

Run on: October 12, 2002, 20:45:36 ; Search time 19.91 Seconds

(without alignments)
48.262 Million cell updates/sec

Title: US-09-408-578a-1

Perfect score: 52

Sequence: 1 PVVAESPCKP 10

Scoring table: BIOSUM62
Gapop 10.0, Gapext 0.5

Searched: 28338 segs, 96089334 residues

Total number of hits satisfying chosen parameters: 11821

Minimum DB seq length: 0
Maximum DB seq length: 50

Post-processing: Minimum Match 0%
Maximum Match 100%

Listing first 45 summaries

Database :
1: PIR1:*
2: PIR2:*
3: PIR3:*
4: PIR4:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	27	51.9	15	2	PS0251
2	26	50.0	14	2	PN0666
3	26	50.0	32	2	S01810
4	26	50.0	47	2	E47395
5	26	50.0	47	2	S14064
6	25	48.1	20	2	I53671
7	25	48.1	30	2	S59482
8	25	48.1	36	2	A82208
9	25	48.1	37	1	R5B536
10	25	48.1	37	2	B32307
11	25	48.1	37	2	T44407
12	25	48.1	37	2	E97282
13	25	48.1	37	2	F90019
14	25	48.1	37	2	AH1776
15	25	48.1	37	2	AI1400
16	25	48.1	39	2	S16978
17	25	48.1	40	2	A28938
18	25	48.1	49	2	D88533
19	25	48.1	50	2	C88533
20	24	46.2	12	2	B39690
21	24	46.2	17	2	S6198
22	24	46.2	27	2	A30323
23	24	46.2	37	2	E81254
24	24	46.2	37	2	G95010
25	24	46.2	39	2	T42897
26	24	46.2	41	2	S10263
27	24	46.2	42	2	T36992
28	24	46.2	45	2	S04941
29	24	46.2	45	2	S10544

30	24	46.2	45	2	S10545	protamine phi-3.3
31	24	46.2	48	2	I61693	myosin - human (fr
32	24	46.2	49	2	PC2062	dihemic cytochrome
33	24	46.2	49	2	A81605	hypothetical prote
34	24	46.2	50	2	D90706	hypothetical prote
35	24	46.2	50	2	F82409	hypothetical prote
36	23	44.2	11	2	P00231	beta-glucosidase (
37	23	44.2	15	2	A54397	ubiquitin-carrier
38	23	44.2	26	2	S06675	apidaecin Ib precu
39	23	44.2	27	2	S71302	IC16 protein - Par
40	23	44.2	30	2	S72626	Small-cell-variant
41	23	44.2	31	2	S50903	fatty acid beta-ox
42	23	44.2	31	2	T36103	hypothetical prote
43	23	44.2	32	2	C61491	seed protein ws-3
44	23	44.2	34	2	S17644	alcohol dehydrogen
45	23	44.2	36	2	A69287	hypothetical prote

ALIGNMENTS

RESULT 1
PS0251
15K protein-5106 - rice (strain Nihonbare) (fragment)
C:Species: Oryza sativa (rice)
C:Date: 03-Feb-1994 #sequence_revision 03-Feb-1994 #text_change 11-Apr-1995
C:Accession: PS0251
R:Tsuigita, A.; Kano, M.
submitted to JIPID, April 1993
A:Reference number: PS0209
A:Accession: PS0251
A:Molecule type: protein
A:Residues: 1-15 <TSU>
A:Experimental source: germ, strain Nihonbare
C:Comment: molecular weight 15K, pI 9.2.

Query Match 51.9%; Score 27; DB 2; Length 15;
Best Local Similarity 33.3%; Pred. No. 1.1e+02;
Matches 3; Conservative 5; Mismatches 1; Indels 0; Gaps 0;

OY 1 PVVAESPCKP 9
DB 5 PIMADXPKE 13

RESULT 2
PN0666
dystrophin-associated glycoprotein A3a-V - rabbit (fragment)
C:Species: Oryctolagus cuniculus (domestic rabbit)
C:Date: 19-May-1994 #sequence_revision 19-May-1994 #text_change 07-May-1999
C:Accession: PN0666
R:Yoshida, M.; Mizuno, Y.; Nonaka, I.; Ozawa, E.
J. Biochem. 114, 634-639, 1993
A:Title: A dystrophin-associated glycoprotein, A3a (one of 43DAG doublets), is retain
A:Reference-number: PN0662; MUID:94156881
A:Accession: PN0666
A:Molecule type: protein
A:Residues: 1-14 <YOS>
C:Comment: This protein is retained in Duchenne type muscular dystrophy muscle.
C:Keywords: glycoprotein; skeletal muscle

Query Match 50.0%; Score 26; DB 2; Length 14;
Best Local Similarity 40.0%; Pred. No. 1.3e+02;
Matches 4; Conservative 2; Mismatches 4; Indels 0; Gaps 0;

OY 1 PVVAESPCKP 10
DB 3 PUMKXPXAP 12

RESULT 3

501810
hemoglobin AIV - tube worm (Lamellibrachia sp.) (fragment)
C:Species: Lamellibrachia sp.
C:Date: 07-Sep-1990 #sequence_revision 07-Sep-1990 #text_change 18-Jun-1993
C:Accession: 501810
R:Suzuki, T.; Takagi, T.; Ohta, S.
Biochem. J. 255, 541-545, 1988
A:Title: N-terminal amino acid sequence of the deep-sea tube worm haemoglobin remarkably
A:Reference number: 501807; MUID:09076216
A:Accession: 501810
A:Molecule type: protein
A:Residues: 1-32 <SUZ>

Query Match 50.0%; Score 26; DB 2; Length 32;
Best Local Similarity 83.3%; Pred. No. 3.6e+02;
Matches 5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

OY 3 VAESP 8
| | | | |
DB 4 VAEPK 9

RESULT 4
E47395
histone H1 H1-1 (clone L92) - midge (Chironomus thummi piger) (fragment)
C:Species: Chironomus thummi piger
C:Date: 24-Feb-1994 #sequence_revision 18-Nov-1994 #text_change 03-May-1996
C:Accession: E47395
R:Schulze, E.; Trieschmann, L.; Schulze, B.; Schmidt, E.R.; Pitzel, S.; Zechel, K.; Gros
Proc. Natl. Acad. Sci. U.S.A. 90, 2481-2485, 1993
A:Title: Structural and functional differences between histone H1 sequence variants with
A:Reference number: A47395; MUID:93211985
A:Accession: E47395
A:Status: preliminary; not compared with conceptual translation
A:Molecule type: DNA
A:Residues: 1-47 <SCH>
A:Note: Sequence extracted from NCBI backbone (NCBIP:128540)
C:Superfamily: histone H1

Query Match 50.0%; Score 26; DB 2; Length 47;
Best Local Similarity 50.0%; Pred. No. 5.3e+02;
Matches 5; Conservative 1; Mismatches 4; Indels 0; Gaps 0;

OY 1 PVVESP 10
| | | | |
DB 10 PAKAKKEKP 19

RESULT 5
S14064
hypothetical protein 2 - Streptomyces griseus (fragment)
C:Species: Streptomyces griseus
C:Date: 19-Mar-1997 #sequence_revision 26-Feb-1998 #text_change 26-Feb-1998
C:Accession: S14064
R:Yigal, T.; Gill, J.A.; Daza, A.; Garcia-Gonzalez, M.D.; Martin, J.F.
Mol. Gen. Genet. 225, 278-288, 1991
A:Title: Cloning, characterization and expression of an alpha-amylase gene from Streptom
A:Reference number: S14062; MUID:9117128
A:Accession: S14064
A:Molecule type: DNA
A:Residues: 1-47 <YIG>
A:Experimental source: strain IMR3570

Query Match 50.0%; Score 26; DB 2; Length 47;
Best Local Similarity 40.0%; Pred. No. 5.3e+02;
Matches 4; Conservative 2; Mismatches 4; Indels 0; Gaps 0;

OY 1 PVVESP 10
| | | | |
DB 22 PAAADTPDAP 31

RESULT 6
I53671
neurofilament heavy subunit - human (fragment)
C:Species: Homo sapiens (man)
C:Date: 02-Jul-1996 #sequence_revision 02-Jul-1996 #text_change 05-Nov-1999
C:Accession: I53671
R:Figlewicz, D.A.; Rouleau, G.A.; Kitzus, A.; Julien, J.P.
Gene 132, 297-300, 1993
A:Title: Polymorphism in the multi-phosphorylation domain of the human neurofilament
A:Reference number: I53671; MUID:94040777
A:Accession: I53671
A:Status: preliminary; translated from GB/EMBL/DDat
A:Molecule type: mRNA
A:Residues: 1-20 <RES>
A:Cross-references: GB:S66488; NID:9452861; PIDN:AAB28609.1; PID:9452862
C:Genetics:
A:Gene: GDB:NEFH
A:Cross-references: GDB:120225; OMIM:162230
A:Map position: 22q12.1-22q13.1

Query Match 48.1%; Score 25; DB 2; Length 20;
Best Local Similarity 55.6%; Pred. No. 3.4e+02;
Matches 5; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

OY 1 PVVESP 9
| | | | |
DB 2 PEKAKSPK 10

RESULT 7
S59482
hydroxyproline-rich cell wall glycoprotein, 136k, major component - kidney bean (fra
N:Alternate names: extensin-like protein
C:Species: Phaseolus vulgaris (kidney bean)
C:Date: 27-Apr-1996 #sequence_revision 19-Jul-1996 #text_change 05-Dec-1998
C:Accession: S59482
R:Wojtaszek, P.; Trechow, J.; Bolwell, G.P.
Plant Mol. Biol. 28, 1075-1087, 1995
A:Title: Specificity in the immobilisation of cell wall proteins in response to diff
A:Reference number: S59481; MUID:96011753
A:Accession: S59482
A:Molecule type: protein
A:Residues: 1-30 <WOJ>
C:Keywords: glycoprotein; hydroxyproline
F:8,9,10,11,12,17,18,19,20,26,27,28,29/Modified site: hydroxyproline (Pro) #status e

Query Match 48.1%; Score 25; DB 2; Length 30;
Best Local Similarity 50.0%; Pred. No. 5.1e+02;
Matches 5; Conservative 0; Mismatches 5; Indels 0; Gaps 0;

OY 1 PVVESP 10
| | | | |
DB 11 PPVWSSPEP 20

RESULT 8
A82208
hypothetical protein VCJ385 [imported] - Vibrio cholerae (strain N16961 serogroup O1
C:Species: Vibrio cholerae
C:Date: 18-Aug-2000 #sequence_revision 20-Aug-2000 #text_change 02-Feb-2001
C:Accession: A82208
R:Heidelberg, J.F.; Eisen, J.A.; Nelson, W.C.; Clayton, R.A.; Gwinn, M.L.; Dodson, R
chardson, D.; Ermolaeva, M.D.; Vamathevan, J.; Bass, S.; Qin, H.; Dragol, I.; Sella
l, R.R.; Mekalanos, J.J.; Venter, J.C.; Fraser, C.M.
Nature 406, 477-483, 2000
A:Title: DNA Sequence of both chromosomes of the cholera pathogen Vibrio cholerae.
A:Reference number: A82035; MUID:20405833
A:Accession: A82208
A:Status: preliminary

A:Molecule type: DNA
 A:Residues: 1-36 <HEI>
 A:Cross-references: GB:AE004217; GB:AE003852; NID:99655866; PIDN:AAF94543.1; GSPDB:GN001
 A:Experimental source: serogroup O1; strain N16961; biotype El TOR
 C:Genetics:
 A:Gene: VC1385
 A:Map position: 1

Query Match 48.1%; Score 25; DB 2; Length 36;
 Best Local Similarity 71.4%; Pred. No. 6.2e+02;
 Matches 5; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Oy 4 AESPCKP 10
 1 1 1 1 1
 Db 10 APEPKAP 16

RESULT 9

R5BS36
 ribosomal protein L36 - Bacillus stearothermophilus
 N:Alternate names: ribosomal protein BL38; ribosomal protein II
 C:Species: Bacillus stearothermophilus
 C>Date: 31-Mar-1991 #sequence_revision 31-Mar-1991 #text_change 07-May-1999
 C:Accession: S08566; S59066
 R:Tanaka, I.; Kimura, M.; Kimura, J.; Dijk, J.
 FEBS Lett. 166, 343-346, 1984

A:Title: The amino acid sequence of two small ribosomal proteins from Bacillus stearothermophilus
 A:Reference number: S07236; MUID:84108949
 A:Accession: S08566
 A:Molecule type: protein
 A:Residues: 1-37 <TAN>

R:Urbard, H.; Kruff, V.; Bischof, O.; Mueller, E.C.; Wittmann-Liebold, B.
 EMBO J. 14, 4578-4586, 1995
 A:Title: Protein-RNA binding features and their structural and functional implications
 A:Reference number: S59051; MUID:96003638
 A:Accession: S59066
 A:Molecule type: protein

A:Residues: 14-31 <URL>
 C:Superfamily: Escherichia coli ribosomal protein L36
 C:Keywords: protein biosynthesis; ribosome
 F:1-37/Product: ribosomal protein L36a #status predicted <MAT>

Query Match 48.1%; Score 25; DB 1; Length 37;
 Best Local Similarity 57.1%; Pred. No. 6.4e+02;
 Matches 4; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

Oy 2 VVAESPK 8
 1 1 1 1 1
 Db 25 VICENPK 31

RESULT 10

B32307
 ribosomal protein L36 - Bacillus subtilis
 N:Alternate names: ribosomal protein B (rpmJ)
 C:Species: Bacillus subtilis
 C>Date: 31-Mar-1992 #sequence_revision 31-Mar-1992 #text_change 20-Jun-2000
 C:Accession: A32307; D69698; B32307
 R:Boylan, S.A.; Suh, J.W.; Thomas, S.M.; Price, C.W.
 J. Bacteriol. 171, 2553-2562, 1989

A:Title: Gene encoding the alpha core subunit of Bacillus subtilis RNA polymerase is cotranscribed with the beta subunit
 A:Reference number: A32307; MUID:89213940
 A:Accession: A32307
 A:Status: not compared with conceptual translation
 A:Molecule type: DNA
 A:Residues: 1-37 <BOY>

A:Cross-references: GB:M2614; NID:9142458; PIDN:AAA22214.1; PID:9142460
 R:Kunst, F.; Ogasawara, N.; Moszer, I.; Albertini, A.M.; Alloni, G.; Azevedo, V.; Berta, C.; Bron, S.; Brouillet, S.; Brusch, C.V.; Capuano, V.; Carter, N.M.; Chd A.; Ehrlich, S.D.; Emmerison, P.T.; Entian, K.D.; Errington, J.; Fabret, C.; Ferrari, E. Nature 350, 249-256, 1997

A:Authors: Foulger, D.; Fritz, C.; Fujita, M.; Fujita, Y.; Funo, S.; Galizzi, A.; Galich, J.; Harwood, C.R.; Henuit, A.; Hilbert, H.; Holsappel, S.; Hosono, S.; Hullo, K.; Koster, P.; Koningsstein, G.; Krogh, S.; Kumano, M.; Kurita, K.; Lapidus, A.; Lardin, A.; Authors: Lauber, J.; Lazarevic, V.; Lee, S.M.; Levine, A.; Liu, H.; Masuda, S.; Ma Y. M.; Ogawa, K.; Ogilwara, A.; Oudga, B.; Park, S.H.; Parro, V.; Pohl, T.M.; Porter, R.; Rieger, M.; Rivolta, C.; Rocha, E.; Roche, B.; Rose, M.; Sadale, Y.; Sato, T.; Scan A:Authors: Schleich, S.; Schwoerer, R.; Scifone, F.; Sekiguchi, J.; Sekowska, A.; S T.; Winters, P.; Wipat, A.; Yamamoto, H.; Yamane, K.; Yasumoto, K.; Yata, K.; Yoshida A:Authors: Yoshikawa, H.F.; Zumbstein, E.; Yoshikawa, H.; Danchin, A.
 A:Title: The complete genome sequence of the Gram-positive bacterium Bacillus subtilis
 A:Reference number: A69580; MUID:96044033
 A:Accession: D69698
 A:Status: nucleic acid sequence not shown; translation not shown
 A:Molecule type: DNA
 A:Residues: 1-37 <KUN>

A:Cross-references: GB:299104; GB:AL009126; NID:92632267; PIDN:CAB11916.1; PID:92632
 A:Experimental source: strain 168
 C:Genetics:
 A:Gene: rpmJ
 C:Superfamily: Escherichia coli ribosomal protein L36
 C:Keywords: protein biosynthesis; ribosome

Query Match 48.1%; Score 25; DB 2; Length 37;
 Best Local Similarity 57.1%; Pred. No. 6.4e+02;
 Matches 4; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

Oy 2 VVAESPK 8
 1 1 1 1 1
 Db 25 VICENPK 31

RESULT 11

T44407
 ribosomal protein L36 rpmJ [imported] - Bacillus halodurans (strain C-125)
 C:Species: Bacillus halodurans
 C>Date: 31-Jan-2000 #sequence_revision 31-Jan-2000 #text_change 15-Jun-2001
 C:Accession: T44407; G83669
 R:Takami, H.; Nakasone, K.; Takaki, Y.; Maeno, G.; Sasaki, R.; Masui, N.; Fujii, F.; I
 Biosci. Biotechnol. Biochem. 63, 452-455, 1999

A:Title: Sequence analysis of a 32-kb region including the major ribosomal protein gene
 A:Reference number: Z22756; MUID:99209008
 A:Accession: T44407
 A:Status: preliminary; translated from GB/EMBL/DBJ
 A:Molecule type: DNA
 A:Residues: 1-37 <TAK>

A:Cross-references: EMBL:AB017508; NID:94512395; PIDN:BAA75295.1; PID:94512428
 A:Experimental source: strain C-125
 R:Takami, H.; Nakasone, K.; Takaki, Y.; Maeno, G.; Sasaki, R.; Masui, N.; Fujii, F.; I
 Nucleic Acids Res. 28, 4317-4331, 2000

A:Title: Complete genome sequence of the alkaliphilic bacterium Bacillus halodurans
 A:Reference number: A83650; MUID:20512582; PMID:11058132
 A:Accession: G83669
 A:Status: preliminary
 A:Molecule type: DNA
 A:Residues: 1-37 <STO>

A:Cross-references: GB:AP001507; GB:BA000004; NID:910172612; PIDN:BA003878.1; GSPDB:G
 C:Genetics:
 A:Gene: rpmJ
 C:Superfamily: Escherichia coli ribosomal protein L36

Query Match 48.1%; Score 25; DB 2; Length 37;
 Best Local Similarity 57.1%; Pred. No. 6.4e+02;
 Matches 4; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

Oy 2 VVAESPK 8
 1 1 1 1 1
 Db 25 VICENPK 31

RESULT 12
E97282
ribosomal protein L36 [imported] - Clostridium acetobutylicum
C.Species: Clostridium acetobutylicum
C.Date: 14-Sep-2001 #sequence_revision 14-Sep-2001 #text_change 14-Sep-2001
C.Accession: E97282
R.Nolling, J.; Breton, G.; Omelchenko, M.V.; Markarova, K.S.; Zeng, Q.; Gibson, R.; Lee, J.; Daly, M.J.; Bennett, G.N.; Koonin, E.V.; Smith, D.R.
J. Bacteriol. 183, 4823-4838, 2001
A.Title: Genome Sequence and Comparative Analysis of the Solvent-Producing Bacterium CLO
A.Reference number: A96900; MUID:21359325; PMID:21359325
A.Accession: E97282
A.Status: Preliminary
A.Molecule type: DNA
A.Residues: 1-37 <KUR>
A.Cross-References: GB:AE001437; PIDN:AAK1048.1; PID:g15026175; GSPDB:GN00168
A.Experimental source: Clostridium acetobutylicum ATCC824
C.Genetics:
A.Gene: CAC3108

Query Match 48.1%; Score 25; DB 2; Length 37;
Best Local Similarity 57.1%; Pred. No. 6.4e+02;
Matches 4; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

OY 2 VVAESPK 8
1: 1:11
DB 25 VICENPK 31

RESULT 13
F90019
50S ribosomal protein L36 [imported] - Staphylococcus aureus (strain N315)
C.Species: Staphylococcus aureus
C.Date: 10-May-2001 #sequence_revision 10-May-2001 #text_change 22-Oct-2001
C.Accession: F90019
R.Kuroda, M.; Ohta, T.; Uchiyama, I.; Baba, T.; Yuzawa, H.; Kobayashi, I.; Cui, L.; Oguc, M.; Mizutani-U, Y.; Kobayashi, N.; Sawano, T.; Inoue, R.; Kaito, C.; Sekimizu, K.; C.; Shiba, T.; Hattori, M.; Ogasawara, N.; Hayashi, H.; Hiramoto, K.
Lancet 357, 1225-1240, 2001
A.Title: Whole genome sequencing of methicillin-resistant Staphylococcus aureus.
A.Reference number: A9758; MUID:21311952; PMID:11418146
A.Accession: F90019
A.Status: Preliminary
A.Molecule type: DNA
A.Residues: 1-37 <KUR>
A.Cross-References: GB:BA000018; PID:g13702027; PIDN:BAK43319.1; GSPDB:GN00149
A.Experimental source: strain N315
C.Genetics:
A.Gene: rpmJ

Query Match 48.1%; Score 25; DB 2; Length 37;
Best Local Similarity 57.1%; Pred. No. 6.4e+02;
Matches 4; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

OY 2 VVAESPK 8
1: 1:11
DB 25 VICENPK 31

RESULT 14
AH1776
ribosomal protein L36 [imported] - Listeria innocua (strain C1p11262)
C.Species: Listeria innocua
C.Date: 27-Nov-2001 #sequence_revision 27-Nov-2001 #text_change 27-Nov-2001
C.Accession: AH1776
R.Glasner, P.; Frangoul, L.; Buchrieser, C.; Amend, A.; Baquero, F.; Berche, P.; Biocker, J.; Dominguez-Bernal, G.; Duchaud, E.; Durand, L.; Dussurget, O.; Entian, K.D.; Fsihl, H.; Jones, L.M.; Karst, U.
Science 294, 849-852, 2001
A.Authors: Kretz, J.; Kuhn, M.; Kunst, F.; Kurapkut, G.; Madueno, E.; Maitournam, A.; Maok, C.; Schlueter, T.; Simoes, N.; Tierrez, A.; Vazquez-Boland, J.A.; Voss, H.; Wehlend,

A.Title: Comparative genomics of Listeria species.
A.Reference number: AB1077; MUID:21537279; PMID:11679669
A.Accession: AH1776
A.Status: Preliminary
A.Molecule type: DNA
A.Residues: 1-37 <GLA>
A.Cross-References: GB:AL592022; PIDN:CAC97984.1; PID:g16415294; GSPDB:GN00178
A.Experimental source: strain C1p11262
C.Genetics:
A.Gene: rpmJ

Query Match 48.1%; Score 25; DB 2; Length 37;
Best Local Similarity 57.1%; Pred. No. 6.4e+02;
Matches 4; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

OY 2 VVAESPK 8
1: 1:11
DB 25 VICENPK 31

RESULT 15
A11400
ribosomal protein L36 [imported] - Listeria monocytogenes (strain EGD-e)
C.Species: Listeria monocytogenes
C.Date: 27-Nov-2001 #sequence_revision 27-Nov-2001 #text_change 27-Nov-2001
C.Accession: A11400
R.Glasner, P.; Frangoul, L.; Buchrieser, C.; Amend, A.; Baquero, F.; Berche, P.; Biocker, J.; Dominguez-Bernal, G.; Duchaud, E.; Durand, L.; Dussurget, O.; Entian, K.D.; Fsihl, H.; Jones, L.M.; Karst, U.
Science 294, 849-852, 2001
A.Authors: Kretz, J.; Kuhn, M.; Kunst, F.; Kurapkut, G.; Madueno, E.; Maitournam, A.; Maok, C.; Schlueter, T.; Simoes, N.; Tierrez, A.; Vazquez-Boland, J.A.; Voss, H.; Wehlend, J.
A.Title: Comparative genomics of Listeria species.
A.Reference number: AB1077; MUID:21537279; PMID:11679669
A.Accession: A11400
A.Status: Preliminary
A.Molecule type: DNA
A.Residues: 1-37 <GLA>
A.Cross-References: GB:NC_003210; PIDN:CAD00687.1; PID:g16412097; GSPDB:GN00177
A.Experimental source: strain EGD-e
C.Genetics:
A.Gene: rpmJ

Query Match 48.1%; Score 25; DB 2; Length 37;
Best Local Similarity 57.1%; Pred. No. 6.4e+02;
Matches 4; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

OY 2 VVAESPK 8
1: 1:11
DB 25 VICENPK 31

Search completed: October 12, 2002, 20:49:44
Job time: 248 sec

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OM protein - protein search, using sw model

Run on: October 12, 2002, 20:48:06 ; Search time 9.8 Seconds
(without alignments)
39.510 Million cell updates/sec

Title: US-09-408-578A-1
Perfect score: 52
Sequence: 1 PVAESPCKP 10

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 105224 seqs, 38719550 residues

Total number of hits satisfying chosen parameters: 3667

Minimum DB seq length: 0
Maximum DB seq length: 50

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database: SWISSPROT_40:*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	26	50.0	32	1	GLB4_LAMSP
2	25	48.1	29	1	TAT_HV123
3	25	48.1	33	1	RL4_HALCU
4	25	48.1	37	1	RL36_BACD
5	25	48.1	37	1	RL36_BACST
6	25	48.1	37	1	RL36_BACSU
7	25	48.1	37	1	RL36_THETH
8	25	48.1	39	1	PSAX_ANAVA
9	25	48.1	49	1	PK06_CAEUL
10	25	48.1	50	1	PK02_CAEUL
11	25	48.1	50	1	PK05_CAEUL
12	25	46.2	16	1	H5_COTJA
13	24	46.2	37	1	CHCD_ANTPO
14	24	46.2	37	1	RL36_CAMPE
15	24	46.2	45	1	H32_TETRO
16	24	46.2	40	1	PH13_MYCA
17	24	46.2	50	1	HOKE_ECOLI
18	23	44.2	17	1	TRP2_LEUMA
19	23	44.2	24	1	CR16_LITXA
20	23	44.2	24	1	CR17_LITXA
21	23	44.2	24	1	CR18_LITXA
22	23	44.2	24	1	CR19_LITXA
23	23	44.2	25	1	CR12_LITCE
24	23	44.2	25	1	CR13_LITCE
25	23	44.2	25	1	CR14_LITCE
26	23	44.2	25	1	CR15_LITCE
27	23	44.2	25	1	CR15_LITCE
28	23	44.2	25	1	CR15_LITCE
29	23	44.2	36	1	Y297_ARCFU
30	23	44.2	39	1	GVPK_SPICC
31	23	44.2	44	1	BABA_BABBO
32	22	42.3	17	1	APID_BOWPA
33	22	42.3	34	1	HIS_STRPU

34	22	42.3	37	1	RK36_CHUV
35	22	42.3	41	1	LAMA_EMENT
36	21	40.4	17	1	SRY_URSAR
37	21	40.4	20	1	HELT_HELHO
38	21	40.4	20	1	ML17_BOVIN
39	21	40.4	31	1	CUS4_LOCMT
40	21	40.4	33	1	PSBT_MAIZE
41	21	40.4	34	1	EMI_ENSMI
42	21	40.4	34	1	PSBT_TOBAC
43	21	40.4	35	1	COPR_CANFA
44	21	40.4	35	1	PSBT_MARPO
45	21	40.4	35	1	PSBT_ORYSA

ALIGNMENTS

RESULT 1	GLB4_LAMSP	STANDARD;	PRT;	32 AA.
ID	P20413;			
AC	01-FEB-1991 (Rel. 17, Created)			
DT	01-FEB-1991 (Rel. 17, Last sequence update)			
DE	01-FEB-1991 (Rel. 17, Last annotation update)			
OS	Giant Hemoglobin AIV chain (Fragment).			
OC	Lamellibrachia sp. (Deep-sea giant tube worm).			
OX	Eukaryota; Metazoa; Vestimentifera; Basibranchia; Lamellibrachida; Lamellibrachidae; Lamellibrachia.			
RN	NCBI_TaxID=6424;			
RP	[1]			
RX	SEQUENCE.			
RA	MEDLINE=89076216; PubMed=3202832;			
RT	Suzuki T., Takagi T., Ohta S.;			
RL	"N-terminal amino acid sequence of the deep-sea tube worm haemoglobin			
CC	Biochem. J. 255:541-545(1988)."			
CC	-1 SUBUNIT: GIANT HEMOGLOBIN IS COMPOSED OF FOUR HEME-CONTAINING			
DR	PIR; S01810; S01810.			
DR	InterPro; IPR000971; Globin.			
DR	PROSITE; PS01033; GLOBIN; PARTIAL.			
KW	Heme; Oxygen transport; transport.			
FT	NON_TER			
FT	32			
FT	SEQUENCE 32 AA; 3695 MW; FD7E1ADABD35FD13 CRC64;			
QY	3 VAEAPK 8			
QY				
DB	4 VAEAPK 9			
Query Match	50.0%; Score 26; DB 1; Length 32;			
Best Local Similarity	83.3%; Pred. No. 1.9e+02;			
Matches	5; Conservative 1; Mismatches 0; Indels 0; Gaps 0;			
RESULT 2	TAT_HV123	STANDARD;	PRT;	29 AA.
ID	TAT_HV123			
AC	P12510;			
DT	01-OCT-1989 (Rel. 12, Created)			
DE	01-OCT-1989 (Rel. 12, Last sequence update)			
DE	16-OCT-2001 (Rel. 40, Last annotation update)			
OS	TAT protein (Transactivating regulatory protein) (Fragment).			
OC	Human immunodeficiency virus type 1 (Zaire 3 isolate) (HIV-1).			
OX	Vitruvius; Retroviral viruses; Retroviridae; Lentivirus.			
RN	NCBI_TaxID=11680;			
RP	[1]			
RX	SEQUENCE FROM N.A.			
RA	MEDLINE=86259728; PubMed=3014529;			
RA	Willey R.W., Rutledge R.A., Dias S., Folks T., Theodore T.,			
RA	Buckler C.E., Martin M.A.;			
RT	"Identification of conserved and divergent domains within the			

RT envelope gene of the acquired immunodeficiency syndrome retrovirus."
 RL Proc. Natl. Acad. Sci. U.S.A. 83:5038-5042(1986)
 CC -1- FUNCTION: TRANSCRIPTIONAL REGULATOR THAT ACTS BY BINDING TO THE
 CC TRANS-ACTIVATING RESPONSIVE SEQUENCE (TAR) RNA ELEMENT AND
 CC ACTIVATES TRANSCRIPTION INITIATION AND/OR ELONGATION FROM THE LTR
 CC PROMOTER.
 CC -1- SUBUNIT: BINDS CYCLIN T1 (BY SIMILARITY).
 CC -1- SUBCELLULAR LOCATION: NUCLEAR; NUCLEOLAR.
 CC -----
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 CC or send an email to license@isb-sib.ch).
 CC -----
 CC EMBL: K03347; AAA5374.1; -
 DR HIV; K03347; TAT523.
 KM Transcription regulation; Activator; RNA-binding; Nuclear protein;
 KW AIDS.
 FT NON_TER 1 1
 SQ SEQUENCE 29 AA; 3229 MW; A8C8E45E70FC192 CRC64;

Query Match 48.1%; Score 25; DB 1; Length 29;
 Best Local Similarity 55.6%; Pred. No. 2.6e+02;
 Matches 5; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

OY 1 PVVAESPK 9
 Db 9 PTGGEPEK 17

RESULT 3
 RL HALCU STANDARD; PRT; 33 AA.
 ID P05967;
 AC 01-NOV-1988 (Rel. 09, Created)
 DT 01-NOV-1988 (Rel. 09, Last sequence update)
 DE 30-MAY-2000 (Rel. 39, Last annotation update)
 DE 50S ribosomal protein L4E (HL9) (Fragment).
 GN RPL4E.
 OS Halobacterium cutirubrum.
 OC Archaea; Euryarchaeota; Halobacteriales; Halobacteriaceae;
 OC Halobacterium.
 OX NCBI_TaxID=2242;
 RN [1]
 RP SEQUENCE.
 RX MEDLINE=84282108; PubMed=6467081;
 RA Matheson A.T., Yaguchi M., Christensen P., Rollin C.F., Hasnain S.;
 RT "Purification, properties, and N-terminal amino acid sequence of
 RT certain 50S ribosomal subunit proteins from the archaeobacterium
 RT Halobacterium cutirubrum."
 RL Can. J. Biochem. Cell Biol. 62:426-433(1984).
 CC -1- SIMILARITY: BELONGS TO THE L4E FAMILY OF RIBOSOMAL PROTEINS.
 DR PIR: S08551; S08551.
 DR InterPro: IPR002136; RIBOSOMAL_L4/L1E.
 DR PROSITE: PS00939; RIBOSOMAL_L1E; PARTIAL.
 KW RIBOSOMAL protein.
 FT NON_TER 33
 SQ SEQUENCE 33 AA; 3753 MW; E0D799DAAF1E3B7B CRC64;

Query Match 48.1%; Score 25; DB 1; Length 33;
 Best Local Similarity 50.0%; Pred. No. 2.9e+02;
 Matches 5; Conservative 1; Mismatches 4; Indels 0; Gaps 0;
 OY 1 PVVAESPK 10
 Db 19 PVEEPEVRP 28

RESULT 4
 RL36_BACSD STANDARD; PRT; 37 AA.
 ID O50631; Q9JPM6;
 AC 15-DEC-1998 (Rel. 37, Created)
 DT 15-DEC-1998 (Rel. 37, Last sequence update)
 DT 16-OCT-2001 (Rel. 40, Last annotation update)
 DE 50S ribosomal protein L36.
 GN RPL36 OR BH0159.
 OS Bacillus halodurans.
 OC Bacteria; Firmicutes; Bacillus/Clostridium group;
 OC Bacillus/Staphylococcus group; Bacillus.
 OX NCBI_TaxID=86655;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=C-125 / JCM 9153;
 RX MEDLINE=99052103; PubMed=9835038;
 RA Nakasone K., Takaki Y., Takami H., Inoue A., Horikoshi K.;
 RT "Cloning and expression of the gene encoding RNA polymerase alpha
 RT subunit from alkaliphilic Bacillus sp. strain C-125."
 RL FEMS Microbiol. Lett. 168:269-276(1998).
 RN [2]
 RP SEQUENCE FROM N.A.
 RC STRAIN=C-125 / JCM 9153;
 RX MEDLINE=99209008; PubMed=10192928;
 RA Takami H., Takaki Y., Nakasone K., Hirama C., Inoue A., Horikoshi K.;
 RT "Sequence analysis of a 32-kb region including the major ribosomal
 RT protein gene clusters from alkaliphilic Bacillus sp. strain C-125."
 RL Biosci. Biotechnol. Biochem. 63:452-455(1999).
 RN [3]
 RP SEQUENCE FROM N.A.
 RC STRAIN=C-125 / JCM 9153;
 RX MEDLINE=20512582; PubMed=11058132;
 RA Takami H., Nakasone K., Takaki Y., Maeno G., Sasaki R., Masui N.,
 RA Fujii F., Hirama C., Nakamura Y., Ogasawara N., Kuhara S.,
 RA Horikoshi K.;
 RT "Complete genome sequence of the alkaliphilic bacterium Bacillus
 RT halodurans and genomic sequence comparison with Bacillus subtilis."
 RL Nucleic Acids Res. 28:4317-4331(2000).
 CC -1- SIMILARITY: BELONGS TO THE L36P FAMILY OF RIBOSOMAL PROTEINS.
 CC -----
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 CC -----
 CC EMBL: AB010082; BAA24191.1; -
 DR EMBL: AB017508; BAA75295.1; -
 DR EMBL: AP001507; BAB03878.1; -
 DR HSSP: P80256; IDPF.
 DR InterPro: IPR000473; RIBOSOMAL_L36.
 DR Pfam: PF00444; RIBOSOMAL_L36; 1.
 DR ProDom: PD002101; RIBOSOMAL_L36; 1.
 DR PROSITE: PS00828; RIBOSOMAL_L36; 1.
 KW RIBOSOMAL protein; Complete proteome.
 FT SEQUENCE 37 AA; 4278 MW; 93A9A8E0714FACF CRC64;

Query Match 48.1%; Score 25; DB 1; Length 37;
 Best Local Similarity 57.1%; Pred. No. 3.2e+02;
 Matches 4; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

OY 2 VVAESPK 8
 Db 25 VICENPK 31

RESULT 5
 RL36_BACST STANDARD; PRT; 37 AA.
 ID RL36_BACST

```

AC P07841;
DT 01-AUG-1988 (Rel. 08, Created)
DT 01-AUG-1988 (Rel. 08, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE 50S ribosomal protein L36 (Ribosomal protein B) (BL38).
DE (BL38).
GN Bactillus stearothermophilus.
OS Bacteria; Firmicutes; Bacillus/Clostridium group;
OC Bacillus/Staphylococcus group; Geobacillus.
OX NCBI_TaxID=1422;
RN [1]
RP MEDLINE=84108949; PubMed=6420194;
RX Tanaka I., Kimura M., Kimura J., Dijk J.;
RT "The amino acid sequence of two small ribosomal proteins from
RL Bactillus stearothermophilus."
FEBS Lett. 166:343-346(1984).
CC -1- SIMILARITY: BELONGS TO THE L36P FAMILY OF RIBOSOMAL PROTEINS.
DR PIR: S08566; R5B836.
DR HSSP: P80256; 1DPE.
DR InterPro: IPR000473; Ribosomal_L36.
DR Pfam: PF00444; Ribosomal_L36; 1.
DR ProDom: PD002101; Ribosomal_L36; 1.
DR PROSITE: PS00828; RIBOSOMAL_L36; 1.
KW Ribosomal protein.
SQ SEQUENCE 37 AA; 4361 MW; 7F6B0920714F4CF7 CRC64;

Query Match 48.1%; Score 25; DB 1; Length 37;
Best Local Similarity 57.1%; Pred. No. 3.2e+02;
Matches 4; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 2 VVAESPK 8
1: 1:11
DB 25 VICENPK 31

RESULT 6
ID RL36_BACSU STANDARD; PRT; 37 AA.
AC P20278;
DT 01-FEB-1991 (Rel. 17, Created)
DT 01-FEB-1991 (Rel. 17, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE 50S ribosomal protein L36 (Ribosomal protein B) (BL38).
DE (BL38).
GN Bactillus subtilis.
OS Bacteria; Firmicutes; Bacillus/Clostridium group;
OC Bacillus/Staphylococcus group; Bacillus.
OX NCBI_TaxID=1423;
RN [1]
RP MEDLINE=89213940; PubMed=2496109;
RX Boylan S.A., Suh J.-W., Thomas S.M., Price C.W.;
RT "Gene encoding the alpha core subunit of Bacillus subtilis RNA
RT polymerase is co-transcribed with the genes for initiation factor 1
RT and ribosomal proteins B, S13, S11, and L17."
J. Bacteriol. 171:2553-2562(1989).
RN [2]
RP SEQUENCE FROM N.A.
RX STRAIN=168 / MARBURG;
MEDLINE=96186897; PubMed=8635744;
RX Suh J.W., Boylan S.A., Oh S.H., Price C.W.;
RT "Genetic and transcriptional organization of the Bacillus subtilis
RT spc-alpha region."
Gene 169:17-23(1996).
RN [3]
RP -1- SIMILARITY: BELONGS TO THE L36P FAMILY OF RIBOSOMAL PROTEINS.
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CC -----
DR EMBL: M26414; AAA22214.1; -
DR EMBL: LA7971; AAB06823.1; -
DR EMBL: Z99104; CAB11916.1; -
DR PIR: B32307; B32307.
DR HSSP: P80256; 1DPE.
DR Subtilist; BG11042; rpmj.
DR InterPro: IPR000473; Ribosomal_L36.
DR Pfam: PF00444; Ribosomal_L36; 1.
DR ProDom: PD002101; Ribosomal_L36; 1.
DR PROSITE: PS00828; RIBOSOMAL_L36; 1.
KW Ribosomal protein; Complete proteome.
SQ SEQUENCE 37 AA; 4305 MW; 7F79A9E0714F4CF7 CRC64;

Query Match 48.1%; Score 25; DB 1; Length 37;
Best Local Similarity 57.1%; Pred. No. 3.2e+02;
Matches 4; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 2 VVAESPK 8
1: 1:11
DB 25 VICENPK 31

RESULT 7
ID RL36_THERM STANDARD; PRT; 37 AA.
AC P80256;
DT 01-OCT-1993 (Rel. 27, Created)
DT 01-OCT-1993 (Rel. 27, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE 50S ribosomal protein L36 (Ribosomal protein B).
GN rpmj OR RPL36.
OS Thermus aquaticus (subsp. thermophilus).
OC Bacteria; Thermus/Delnococcus group; Thermus group; Thermus.
OX NCBI_TaxID=274;
RN [1]
RP SEQUENCE.
RC STRAIN=HB8 / ATCC 27634;
RC Boyesen R.I., Schroeder W., Erdmann V.A.;
RL Submitted (SEP-1993) to the SWISS-PROT data bank.
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=HB8 / ATCC 27634;
MEDLINE=95098837; PubMed=9880810;
RX Wada T., Yamazaki T., Kuramitsu S., Kyogoku Y.;
RT "Cloning of the RNA polymerase alpha subunit gene from Thermus
RT thermophilus HB8 and characterization of the protein."
J. Biochem. 125:143-150(1999).
RN [3]
RP STRUCTURE BY NMR.
RX MEDLINE=20124006; PubMed=10656825;
RX Hard T., Rak A., Allard P., Kloo L., Garber M.;
RT "The solution structure of ribosomal protein L36 from Thermus
RT thermophilus reveals a zinc-ribon-like fold."
J. Mol. Biol. 296:169-180(2000).
RN [4]
RP -1- SIMILARITY: BELONGS TO THE L36P FAMILY OF RIBOSOMAL PROTEINS.
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CC -----
DR EMBL: AB024328; BAA75545.1; -
DR PDB: 1DPE; 16-FEB-00.
DR PDB: 1DG2; 08-DEC-99.

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DR InterPro: IPR000473; Ribosomal_L36.
 DR Pfam: PF00444; Ribosomal_L36; 1.
 DR Prodom: PD002101; Ribosomal_L36; 1.
 DR PROSITE: PS00828; RIBOSOMAL_L36; 1.
 KW Ribosomal protein; 3D-structure.
 SQ SEQUENCE 37 AA; 4421 MW; 439072E1737E2AE8 CRC64;

Query Match 48.1%; Score 25; DB 1; Length 37;
 Best Local Similarity 57.1%; Pred. No. 3.2e+02;
 Matches 4; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 2 VVAESPK 8
 DB 25 VICENPK 31

RESULT 8
 PSAX_ANAVA STANDARD; PRT; 39 AA.
 AC P23319;
 DT 01-NOV-1991 (Rel. 20, Created)
 DT 01-NOV-1991 (Rel. 20, Last sequence update)
 DT 01-MAR-2002 (Rel. 41, Last annotation update)
 DE Photosystem I 4.8 kDa protein (Fragment).
 GN PSAX.
 OS Anabaena variabilis.
 OC Bacteria; Cyanobacteria; Nostocales; Nostocaceae; Anabaena.
 OX NCBI_TaxID=1172;
 RN [1]
 RP SEQUENCE.
 RC STRAIN-PCC 7937 / ATCC 29413;
 RX MEDLINE-91348228; PubMed-1908790;
 RA Ikeuchi M., Nynus K.V., Inoue Y., Pakrasi H.B.;
 RT "Identities of four low-molecular-mass subunits of the photosystem I
 complex from Anabaena variabilis ATCC 29413. Evidence for the
 presence of the psal gene product in a cyanobacterial complex.";
 RL FEBS Lett. 287:5-9(1991)
 CC -1- SIMILARITY: BELONGS TO THE PSAX FAMILY.
 DR PIR; S16978; S16978.
 KW Photosystem I; Photosynthesis.
 FT NON_TER 39
 SQ SEQUENCE 39 AA; 4141 MW; E762CF2381CDECA7 CRC64;

Query Match 48.1%; Score 25; DB 1; Length 39;
 Best Local Similarity 50.0%; Pred. No. 3.4e+02;
 Matches 5; Conservative 1; Mismatches 4; Indels 0; Gaps 0;

QY 1 PVVAESPKP 10
 DB 6 PAVANTGAKP 15

RESULT 9
 YK06_CAEEL STANDARD; PRT; 49 AA.
 AC P34301;
 DT 01-FEB-1994 (Rel. 28, Created)
 DT 01-FEB-1994 (Rel. 28, Last sequence update)
 DT 01-MAR-2002 (Rel. 41, Last annotation update)
 DE Hypothetical 5.0 kDa protein C06E1.6 in chromosome III.
 GN C06E1.6
 OS Caenorhabditis elegans.
 OC Eukaryota; Metazoa; Nematoda; Chromadorea; Rhabditida; Rhabditoidae;
 OC Rhabditidae; Peloderinae; Caenorhabditis.
 OX NCBI_TaxID=6239;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN-BRISTOL N2;
 RX MEDLINE-94150718; PubMed-7906398;
 RA Wilson R., Almscough R., Anderson K., Baynes C., Berks M.,
 RA Bonfield J., Burton J., Connell M., Copsey T., Cooper J., Coulson A.,

RA Craxton M., Dear S., Du Z., Durlin R., Favell A., Fraser A.,
 RA Fulton L., Gardner A., Green P., Hawkins T., Hillier L., Jier M.,
 RA Johnston L., Jones M., Kershaw J., Kirsten J., Laister N.,
 RA Latreille P., Lightning J., Lloyd C., Mortimore B., O'Callaghan M.,
 RA Parsons J., Percy C., Rifken L., Roopie A., Saunders D., Showkeen R.,
 RA Sims M., Smalton N., Smith A., Smith M., Sonhammer E., Staden R.,
 RA Sulston J., Thierry-Mieg J., Thomas K., Vaudin M., Vaughan K.,
 RA Waterston R., Watson A., Weinstock L., Wilkinson-Sproat J.,
 RA Wohlsman P.,
 RT "2.2 Mb of contiguous nucleotide sequence from chromosome III of C.
 elegans.";
 RL Nature 368:32-38(1994).

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DR EMBL; L16559; AAA27937.1;
 DR Wormpep; C06E1.6; CE00061.
 KW Hypothetical protein.
 SQ SEQUENCE 49 AA; 4970 MW; 8FB975FB9E1AA49B CRC64;

Query Match 48.1%; Score 25; DB 1; Length 49;
 Best Local Similarity 83.3%; Pred. No. 4.2e+02;
 Matches 5; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 4 AESPK 9
 DB 44 AERPK 49

RESULT 10
 YK02_CAEEL STANDARD; PRT; 50 AA.
 AC P34297;
 DT 01-FEB-1994 (Rel. 28, Created)
 DT 01-FEB-1994 (Rel. 28, Last sequence update)
 DT 01-MAR-2002 (Rel. 41, Last annotation update)
 DE Hypothetical 5.0 kDa protein C06E1.2 in chromosome III.
 GN C06E1.2
 OS Caenorhabditis elegans.
 OC Eukaryota; Metazoa; Nematoda; Chromadorea; Rhabditida; Rhabditoidae;
 OC Rhabditidae; Peloderinae; Caenorhabditis.
 OX NCBI_TaxID=6239;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN-BRISTOL N2;
 RX MEDLINE-94150718; PubMed-7906398;
 RA Wilson R., Almscough R., Anderson K., Baynes C., Berks M.,
 RA Bonfield J., Burton J., Connell M., Copsey T., Cooper J., Coulson A.,
 RA Craxton M., Dear S., Du Z., Durlin R., Favell A., Fraser A.,
 RA Fulton L., Gardner A., Green P., Hawkins T., Hillier L., Jier M.,
 RA Johnston L., Jones M., Kershaw J., Kirsten J., Laister N.,
 RA Latreille P., Lightning J., Lloyd C., Mortimore B., O'Callaghan M.,
 RA Parsons J., Percy C., Rifken L., Roopie A., Saunders D., Showkeen R.,
 RA Sims M., Smalton N., Smith A., Smith M., Sonhammer E., Staden R.,
 RA Sulston J., Thierry-Mieg J., Thomas K., Vaudin M., Vaughan K.,
 RA Waterston R., Watson A., Weinstock L., Wilkinson-Sproat J.,
 RA Wohlsman P.,
 RT "2.2 Mb of contiguous nucleotide sequence from chromosome III of C.
 elegans.";
 RL Nature 368:32-38(1994).

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DR EMBL: V00080: CAA23422.1: -
 KW Eggshell; Chorion; Repeat; Multigene family.
 FT NON_TER 1 1
 FT DOMAIN 1 1 CENTRAL DOMAIN.
 FT DOMAIN 27 >37 RIGHT ARM.
 FT NON_TER 37 37
 SQ SEQUENCE 37 AA; 3615 MW; 2EF1FF446F4D532C CRC64;

Query Match 46.2%; Score 24; DB 1; Length 37;
 Best Local Similarity 66.7%; Pred. No. 4.8e+02;
 Matches 4; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

OY 2 VVAESP 7
 :||:|
 Db 23 IVAETP 28

RESULT 14
 RL36_CAMTE STANDARD; PRT; 37 AA.
 ID RL36_CAMTE
 AC G9PM84;
 DT 16-OCT-2001 (Rel. 40, Created)
 DT 16-OCT-2001 (Rel. 40, Last sequence update)
 DT 16-OCT-2001 (Rel. 40, Last annotation update)
 DE 50S ribosomal protein L36.
 GN RPMJ OR CUI591.
 OS Campylobacter jejuni.
 OC Bacteria; Proteobacteria; epsilon subdivision; Campylobacter group;
 CC Campylobacter.
 OX NCBI_TaxID:197;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN-NCTC 11168;
 RX MEDLINE-20150912; PubMed-10688204;
 RA Parish J., Wren B.W., Mungall K., Ketley J.M., Churcher C.,
 RA Basham D., Chillingworth T., Davies R.M., Feltham T., Holtroyd S.,
 RA Jørgensen K., Kariyasek A.V., Moule S., Pallen M.J., Penn C.W.,
 RA Quail M.A., Rajandream M.A., Rutherford K.M., van Vliet A.H.M.,
 RA Whitehead S., Barrett B.G.;
 RT "The genome sequence of the food-borne pathogen Campylobacter jejuni
 RT reveals hypervariable sequences."
 RL Nature 403:665-668(2000).
 CC -1- SIMILARITY: BELONGS TO THE L36P FAMILY OF RIBOSOMAL PROTEINS.
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 CC -----
 CC EMBL: AL139079; CAB73579.1;
 DR InterPro: IPR000473; Ribosomal_L36.
 DR Pfam: PF00444; Ribosomal_L36.1.
 DR PROSITE: PS00828; RIBOSOMAL_L36.1.
 KW Ribosomal protein; Complete proteome.
 SQ SEQUENCE 37 AA; 4364 MW; 4548AEC5256D94ED CRC64;

Query Match 46.2%; Score 24; DB 1; Length 37;
 Best Local Similarity 42.9%; Pred. No. 4.8e+02;
 Matches 3; Conservative 3; Mismatches 1; Indels 0; Gaps 0;

OY 2 VVAESP 8
 :||:|
 Db 25 IICENPK 31

RESULT 15
 H32_TETBO STANDARD; PRT; 40 AA.
 ID H32_TETBO

AC P17319;
 DT 01-AUG-1990 (Rel. 15, Created)
 DT 01-AUG-1990 (Rel. 15, Last sequence update)
 DT 15-JUL-1999 (Rel. 38, Last annotation update)
 DE Histone H3.2 (fragment).
 OS Tetrahymena borealis.
 OC Eukaryota; Alveolata; Ciliophora; Oligohymenophorea; Hymenostomatida;
 OC Tetrahymena; Tetrahymena.
 OX NCBI_TaxID=5893;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN-W23;
 RX MEDLINE-90221813; PubMed-2129549;
 RA Brunk C.F., Sadler L.A.;
 RT "Characterization of the promoter region of Tetrahymena genes."
 RL Nucleic Acids Res. 18:323-329(1990).
 RN [2]
 RP SEQUENCE FROM N.A.
 RC STRAIN-W23;
 RX MEDLINE-90219078; PubMed-2129541;
 RA Brunk C.F., Kahn R.W., Sadler L.A.;
 RT "Phylogenetic relationships among Tetrahymena species determined
 RT using the polymerase chain reaction."
 RL J. Mol. Evol. 30:290-297(1990).
 CC -1- FUNCTION: HISTONE H3, ALONG WITH HISTONE H4, PLAYS A CENTRAL ROLE
 CC IN NUCLEOSOME FORMATION.
 CC -1- SUBUNIT: THE NUCLEOSOME IS AN OCTAMER CONTAINING TWO MOLECULES OF
 CC H2A, H2B, H3, AND H4, WHICH WRAP APPROXIMATELY 146 BP OF DNA.
 CC -1- SIMILARITY: BELONGS TO THE HISTONE H3 FAMILY.
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 CC -----
 CC EMBL: X17128; CAA34990.1; ALT_SEQ.
 DR PIR: S10263; S10263.
 DR InterPro: IPR00164; Histone_H3.
 DR PROSITE: PS00322; HISTONE_H3-1; PARTIAL.
 DR PROSITE: PS00959; HISTONE_H3-2; PARTIAL.
 KW Nuclear protein; Chromosomal protein; DNA-binding; Nucleosome core.
 FT INIT MET 0
 FT NON_TER 40 40
 SQ SEQUENCE 40 AA; 4183 MW; E79CED7EBE66EE02 CRC64;

Query Match 46.2%; Score 24; DB 1; Length 40;
 Best Local Similarity 66.7%; Pred. No. 5.2e+02;
 Matches 4; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

OY 4 AESPPK 9
 :||:|
 Db 13 AEAPRK 18

Search completed: October 12, 2002, 20:50:45
 Job time: 159 sec

GenCore version 4.5
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OM protein - protein search, using sw model

Run on: October 12, 2002, 20:47:16 ; Search time 25.85 Seconds

(without alignments)
66.923 Million cell updates/sec

Title: US-09-408-578A-1

Perfect score: 52

Sequence: 1 PVVAESPCKP 10

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 562222 seqs, 172994929 residues

Total number of hits satisfying chosen parameters: 29986

Minimum DB seq length: 0

Maximum DB seq length: 50

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

SPTREMBL.19:**
1: sp_archaea:*
2: sp_bacteria:*
3: sp_fungi:*
4: sp_human:*
5: sp_invertebrate:*
6: sp_mammal:*
7: sp_mhc:*
8: sp_organelle:*
9: sp_phage:*
10: sp_plant:*
11: sp_rodent:*
12: sp_virus:*
13: sp_vertebrate:*
14: sp_unclassified:*
15: sp_virus:*
16: sp_bacteriap:*
17: sp_archaeap:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	30	57.7	22	13	Q9P655 oncorhynchu
2	28	53.8	12	4	Q9UC37 homo sapien
3	28	53.8	41	9	Q9MC40 bacterioph
4	28	53.8	47	16	Q9P230 thizobium m
5	27	51.9	25	7	019452 mus musculu
6	27	51.9	37	11	088771 rattus norv
7	27	51.9	39	2	P82570 streptococ
8	27	51.9	42	5	Q95R01 caenorhadi
9	27	51.9	47	11	Q9JIC8 mus musculu
10	26	50.0	29	8	Q9GB18 melidectes
11	26	50.0	47	12	084028 influenza a
12	25	48.1	18	4	Q96104 homo sapien
13	25	48.1	20	4	Q16070 homo sapien
14	25	48.1	22	10	Q9SLV6 nicotiana t
15	25	48.1	27	4	Q9UMH7 homo sapien
16	25	48.1	29	5	Q95SE3 drosophila

ALIGNMENTS

17	25	48.1	30	3	Q9UR69	Q9UR69 trameles ve
18	25	48.1	32	6	Q9TR01	Q9TR01 bos taurus
19	25	48.1	34	11	Q9C8T3	Q9C8T3 mus musculu
20	25	48.1	36	16	Q9K574	Q9K574 vibrio chol
21	25	48.1	37	16	Q97EK2	Q97EK2 clostridium
22	25	48.1	37	16	Q99S42	Q99S42 staphylococ
23	25	48.1	37	16	Q927N0	Q927N0 listeria in
24	25	48.1	43	4	Q14909	Q14909 homo sapien
25	25	48.1	45	5	Q9VB16	Q9VB16 drosophila
26	25	48.1	48	2	Q9KJKA	Q9KJKA fischerella
27	25	48.1	49	4	Q96G07	Q96G07 homo sapien
28	25	47.1	44	4	Q95537	Q95537 homo sapien
29	24.5	46.2	14	10	Q94IT6	Q94IT6 fragaria nu
30	24	46.2	14	11	Q9OVF3	Q9OVF3 rattus sp.
31	24	46.2	15	10	Q9S8F1	Q9S8F1 zea mays (m
32	24	46.2	18	8	Q9ZT79	Q9ZT79 idris sp. c
33	24	46.2	20	6	Q9TR50	Q9TR50 bos taurus
34	24	46.2	21	2	Q9X3K2	Q9X3K2 prochloroco
35	24	46.2	24	6	Q9BG18	Q9BG18 oryctolagus
36	24	46.2	25	6	Q9BG19	Q9BG19 chetrogaleu
37	24	46.2	28	6	Q9W0N9	Q9W0N9 cercopithe
38	24	46.2	32	4	Q75210	Q75210 homo sapien
39	24	46.2	32	4	Q90PL5	Q90PL5 homo sapien
40	24	46.2	36	5	Q61188	Q61188 colpidium c
41	24	46.2	37	4	Q9TOR7	Q9TOR7 equus cabal
42	24	46.2	37	4	Q15685	Q15685 homo sapien
43	24	46.2	38	2	Q97M62	Q97M62 streptococ
44	24	46.2	38	2	Q9X318	Q9X318 salmonella
45	24	46.2	38	2	Q9X304	Q9X304 salmonella

RESULT 1
Q9P655; PRELIMINARY; PRT; 22 AA.
ID Q9P655; MEDLINE-92082492; PubMed-1747124;
AC Q9P655; DT 01-MAY-2000 (TREMBLrel. 13, Created)
DT 01-MAY-2000 (TREMBLrel. 13, Last sequence update)
DT 01-MAY-2000 (TREMBLrel. 13, Last annotation update)
DE HISTONE H1B.
OS Oncorhynchus mykiss (Rainbow trout) (Salmo gairdneri).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Actinopterygii; Neopterygii; Teleostei; Euteleostei;
OC Protacanthopterygii; Salmoniformes; Salmonidae; Oncorhynchus.
OX NCBI_TaxID-8022;
RN [1]
RP SEQUENCE.
RX MEDLINE-92082492; PubMed-1747124;
RA Davie J.R., Delcuve G.P.;
RT "Characterization and chromatin distribution of the H1 histones and
RT high-mobility-group non-histone chromosomal proteins of trout liver
RT and hepatocellular carcinoma.";
RL Biochem. J. 280:491-497(1991).
SQ SEQUENCE 22 AA; 2132 MW; 3E90388F68189AE3 CRC64;

Query Match 57.7%; Score 30; DB 13; Length 22;
Best Local Similarity 71.4%; Pred. No. 97;
Matches 5; Conservative 2; Mismatches 0; Indels 0; Gaps 0;
QY 4 AESPCKP 10
DB 14 AKAPCKP 20
RESULT 2
ID Q9UC37; PRELIMINARY; PRT; 12 AA.
AC Q9UC37; DT 01-MAY-2000 (TREMBLrel. 13, Created)
DT 01-MAY-2000 (TREMBLrel. 13, Last sequence update)

DT 01-MAY-2000 (TREMBLrel. 13, Last annotation update)
 DE Alpha B CRYSTALLIN FRAGMENT 5.
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
 OX NCBI_TaxID=9606;
 RN [1]
 RP SEQUENCE.
 RX MEDLINE=92218434; PubMed=1560006;
 RA Kato K., Shinohara H., Goto S., Inaguma Y., Morishita R., Asano T.;
 RT "Copurification of small heat shock protein with alpha B crystallin
 from human skeletal muscle.";
 RL J. Biol. Chem. 267:7718-7725(1992).
 SO SEQUENCE 12 AA; 1268 MW; D37BD529CC1B2CD CRC64;

Query Match 53.8%; Score 28; DB 4; Length 12;
 Best Local Similarity 55.6%; Pred. No. 1.2e+02;
 Matches 5; Conservative 1; Mismatches 3; Indels 0; Gaps 0;

OY 1 PVVAESPKK 9
 DB 4 PAVTAPRK 12

RESULT 3
 Q9MC40 PRELIMINARY; PRT; 41 AA.
 AC Q9MC40;
 DT 01-OCT-2000 (TREMBLrel. 15, Created)
 DT 01-OCT-2000 (TREMBLrel. 15, Last sequence update)
 DT 01-DEC-2001 (TREMBLrel. 19, Last annotation update)
 DE ORF92.
 GN ORF92.
 OS Bacteriophage D3.
 OC Viruses; dsDNA viruses, no RNA stage; Caudovirales; Siphoviridae;
 OC Lambda phage group.
 OX NCBI_TaxID=31535;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=20042341; PubMed=10572124;
 RA Glaktian Z.A., Kropinski A.M.;
 RT "Cloning and analysis of the capsid morphogenesis genes of Pseudomonas
 aeruginosa bacteriophage D3; another example of protein chain malty";
 RL J. Bacteriol. 181:7221-7227(1999).
 RN [2]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=20485557; PubMed=11029426;
 RA Kropinski A.M.;
 RT "Sequence of the Genome of the Temperate, Serotype-Converting,
 Pseudomonas aeruginosa Bacteriophage D3.";
 RL J. Bacteriol. 182:6066-6074(2000).
 DR EMBL: AF165214; AAF80768.1;
 SO SEQUENCE 41 AA; 4629 MW; 9632B19C8D142821 CRC64;

Query Match 53.8%; Score 28; DB 9; Length 41;
 Best Local Similarity 62.5%; Pred. No. 4.1e+02;
 Matches 5; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

OY 3 VAESPKK 10
 DB 29 VAIKPKRP 36

RESULT 4
 Q92P30 PRELIMINARY; PRT; 47 AA.
 AC Q92P30;
 DT 01-DEC-2001 (TREMBLrel. 19, Created)
 DT 01-DEC-2001 (TREMBLrel. 19, Last sequence update)
 DT 01-DEC-2001 (TREMBLrel. 19, Last annotation update)
 DE HYPOTHETICAL PROTEIN SMG04310.

GN SMG04310.
 OS Rhizobium meliloti (Sinorhizobium meliloti).
 OC Bacteria; Proteobacteria; alpha subdivision; Rhizobiaceae group;
 OC Rhizobiaceae; Sinorhizobium.
 OX NCBI_TaxID=382;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=1021;
 RX MEDLINE=21368234; PubMed=11474104;
 RA Gallbert F., Finan T.M., Long S.R., Puehler A., Abola P., Ampe F.,
 RA Barloy-Hubler F., Barnett M.J., Becker A., Bolstad P., Botte G.,
 RA Boutry M., Bowser L., Buhrmester J., Cadieu E., Capela D., Chain P.,
 RA Cowie A., Davis R.W., Dreano S., Federspiel N.A., Fisher R.F.,
 RA Gloux S., Godrie T., Goffeau A., Golding B., Guzy J., Gurjal M.,
 RA Hernandez-Lucas T., Hong A., Hulzer L., Hyman R.W., Jones T., Kahn D.,
 RA Kahn M.L., Kalman S., Keating D.H., Kiss E., Komp C., Lelaure V.,
 RA Masny D., Palm C., Peck M.C., Pohl T.M., Portetelle D., Purnelle B.,
 RA Ramsperger U., Surzycki R., Thebault P., Vandenbol M.,
 RA Vorholter F.J., Weidner S., Wells D.H., Wong K., Yeh K.-C., Batut J.;
 RT "The composite genome of the legume symbiont Sinorhizobium meliloti.";
 RL Science 293:668-672(2001).
 DR EMBL: AL591789; CAC46543.1;
 KW Hypothetical protein; Complete proteome.
 SO SEQUENCE 47 AA; 5038 MW; 58EB722379F00CD8 CRC64;

Query Match 53.8%; Score 28; DB 16; Length 47;
 Best Local Similarity 66.7%; Pred. No. 4.7e+02;
 Matches 6; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

OY 2 VVAESPKK 10
 DB 29 VVAEAPVP 37

RESULT 5
 O19452 PRELIMINARY; PRT; 25 AA.
 AC O19452;
 DT 01-JAN-1998 (TREMBLrel. 05, Created)
 DT 01-JAN-1998 (TREMBLrel. 05, Last sequence update)
 DT 01-DEC-2001 (TREMBLrel. 19, Last annotation update)
 DE MHC CLASS II-ASSOCIATED INVARIANT CHAIN (FRAGMENT).
 OS Mus musculus (Mouse).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 OX NCBI_TaxID=10090;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=90257363; PubMed=2111346;
 RA Fades A.-M., Little M., Ramsdorf H.J.;
 RT "The IFN-gamma response of the murine invariant chain gene is mediated
 by a complex enhancer that includes several MHC class II consensus
 elements.";
 RL J. Immunol. 144:4399-4409(1990).
 DR EMBL: M35872; AAA37897.1;
 FT NON-TER 25
 SO SEQUENCE 25 AA; 2973 MW; D22EADA1B6036FCC CRC64;

Query Match 51.9%; Score 27; DB 7; Length 25;
 Best Local Similarity 30.0%; Pred. No. 3.9e+02;
 Matches 3; Conservative 4; Mismatches 3; Indels 0; Gaps 0;

OY 1 PVVAESPKK 10
 DB 15 PILGNRPK 24

RESULT 6
 O88771 PRELIMINARY; PRT; 37 AA.
 ID O88771
 AC O88771;

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DT 01-NOV-1998 (TREMBLrel. 08, Created)
DT 01-NOV-1998 (TREMBLrel. 08, Last sequence update)
DE 01-DEC-2001 (TREMBLrel. 19, Last annotation update)
DE 265 PROTEASOME P112 SUBUNIT (FRAGMENT).
OS Rattus norvegicus (Rat).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
OX NCBI_TaxID=10116;
RN (1)
RP SEQUENCE FROM N.A.
RC TISSUE=BRAIN;
RX MEDLINE=98434650; PubMed=9757068;
RA Seidel B., Kellhoff G., Reinheckel T., Wolf G.;
RT "Differentially expressed genes in hippocampal cell cultures in
RL response to an excitotoxic insult by quinolinic acid.";
RM Mol. Brain Res. 60:296-300(1998).
DR EMBL, AJ007476; CAA07524.1; -.
KW Proteasome.
FT NON_TER 1 1
FT NON_TER 37 37
SQ SEQUENCE 37 AA; 4207 MW; 96B6333E4549A626 CRC64;

Query Match
Best Local Similarity 51.9%; Score 27; DB 11; Length 37;
Matches 5; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

OY 1 PVASPK 8
DB 11 PVASPK 18

RESULT 7
P82570 PRELIMINARY; PRT; 39 AA.
AC P82570;
DT 01-OCT-2000 (TREMBLrel. 15, Created)
DT 01-OCT-2000 (TREMBLrel. 15, Last sequence update)
DT 01-JUN-2001 (TREMBLrel. 17, Last annotation update)
DE HYPOXANTHINE-GUANINE PHOSPHORIBOSYLTRANSFERASE (EC 2.4.2.8) (HGPR)
DE (HGPR) (FRAGMENTS).
OS Streptococcus pyogenes.
OC Bacteria; Firmicutes; Bacilli; Clostridium group; Streptococcaceae;
OC Streptococcus.
OX NCBI_TaxID=1314;
RN (1)
RP SEQUENCE, AND MASS SPECTROMETRY.
RC STRAIN=JRS4;
RA Hogan D.A., Du P., Stevenson T.I., Whitton M., Kilby G.W., Rogers J.,
RA VanBogelen R.A.;
RT "Two-dimensional gel electrophoresis map of Streptococcus pyogenes
RT proteins.";
RL Submitted (MAY-2000) to the SWISS-PROT data bank.
CC -1 CATALYTIC ACTIVITY: IMP + PYROPHOSPHATE = HYPOXANTHINE +
CC 5-PHOSPHO-ALPHA-D-RIBOSE 1-DIPHOSPHATE (GUANINE CAN REPLACE
CC HYPOXANTHINE TO PRODUCE GMP).
CC -1 PATHWAY: PURINE SALVAGE.
CC -1 SUBCELLULAR LOCATION: CYTOPLASMIC (BY SIMILARITY).
CC -1 MASS SPECTROMETRY: MW=35707.93; METHOD=ELECTROSPRAY.
CC -1 SIMILARITY: BELONGS TO THE PURINE/PYRIMIDINE
CC PHOSPHORIBOSYLTRANSFERASE FAMILY.
DR InterPro: IPR002375; Pur_Pyri_Pi_Transf.
DR PROSITE: PS00103; PUR_PYR_Pi_TRANSFER; FALSE_NEG.
KW transferase; Glycosyltransferase; Purine salvage; Magnesium.
FT NON_TER 1 1
FT NON_TER 14 15
FT NON_TER 27 28
FT NON_TER 39 39
SQ SEQUENCE 39 AA; 4167 MW; 0FE33A096B1BCA1F CRC64;

Query Match
Best Local Similarity 51.9%; Score 27; DB 2; Length 39;
Matches 75.0%; Pred. No. 6e+02;

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Matches 6; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

OY 2 VVASPK 9
DB 11 VVASPK 15

RESULT 8
O95R01 PRELIMINARY; PRT; 42 AA.
AC O95R01;
DT 01-DEC-2001 (TREMBLrel. 19, Created)
DT 01-DEC-2001 (TREMBLrel. 19, Last sequence update)
DE 01-DEC-2001 (TREMBLrel. 19, Last annotation update)
DE HYPOPHETICAL 4.7 KDA PROTEIN.
GN K07B1.6
OS Caenorhabditis elegans.
OC Eukaryota; Metazoa; Nematoda; Chromadorea; Rhabditida; Rhabditoidea;
OC Rhabditidae; Peloderinae; Caenorhabditis.
OX NCBI_TaxID=6239;
RN (1)
RP SEQUENCE FROM N.A.
RC STRAIN=BRISTOL N2;
RX MEDLINE=99069613; PubMed=9851916;
RT None;
RT "Genome sequence of the nematode C. elegans: a platform for
RT investigating biology. The C. elegans Sequencing Consortium.";
RT Science 282:2012-2018(1998).
RN (2)
RP SEQUENCE FROM N.A.
RC STRAIN=BRISTOL N2;
RA Pauley A., Gattung S.;
RT "The sequence of C. elegans cosmid K07B1.";
RL Submitted (MAY-1997) to the EMBL/Genbank/DBJ databases.
RN (3)
RP SEQUENCE FROM N.A.
RC STRAIN=BRISTOL N2;
RA Waterston R.;
RT "Direct Submission.";
RL Submitted (OCT-2001) to the EMBL/Genbank/DBJ databases.
DR EMBL, AF003384; AAL06041.1; -.
KW Hypothetical protein.
SQ SEQUENCE 42 AA; 4709 MW; 390F7ADF3BF906E9 CRC64;

Query Match
Best Local Similarity 51.9%; Score 27; DB 5; Length 42;
Matches 4; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

OY 5 ESPKRP 10
DB 11 QSPKRP 16

RESULT 9
O9JIC8 PRELIMINARY; PRT; 47 AA.
AC O9JIC8;
DT 01-OCT-2000 (TREMBLrel. 15, Created)
DT 01-OCT-2000 (TREMBLrel. 15, Last sequence update)
DE 01-DEC-2001 (TREMBLrel. 19, Last annotation update)
DE SMN PROTEIN (FRAGMENT).
GN SMN.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN (1)
RP SEQUENCE FROM N.A.
RC STRAIN=C57BL/6J;
RX MEDLINE=20414734; PubMed=10958634;
RA Growney J.D., Dietrich W.F.;
RT "High-resolution genetic and physical map of the Lgn1 interval in

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RT C57BL/6J implicates Naip2 or Naip5 in Legionella pneumophila
RT pathogenesis."
RL Genome Res. 10:1158-1171(2000).
DR EMBL: AF240503; AAF81197.1; -.
DR MGD: MGI:109257; Smm.
FT NON_TER
FT NON_TER
SQ SEQUENCE 47 AA; 5160 MW; 1CD41D6E32BC7126 CRC64;

Query Match
Best Local Similarity 51.9%; Score 27; DB 11; Length 47;
Matches 4; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

QY 3 VAESPCKP 10
DB 15 ICETPDKP 22

RESULT 10
09GB18 PRELIMINARY; PRT; 29 AA.
AC 09GB18;
DT 01-MAR-2001 (TREMBLrel. 16, Created)
DT 01-MAR-2001 (TREMBLrel. 16, Last sequence update)
DE 01-JUN-2001 (TREMBLrel. 17, Last annotation update)
DE CYTOCHROME C OXIDASE SUBUNIT II (FRAGMENT).
GN COIT.
OS Melidectes belfordi (Belford's melidectes).
OC Mitochondrion.
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Archosauria; Aves; Neognathae; Passeriformes; Meliphagidae;
OC Melidectes.
OX NCBI_TaxID=43161;
RN [1]
RP SEQUENCE FROM N.A.
RA Silkas B., Jones I.B., Derrickson S.R., Fleischner R.C.;
RT "Phylogenetic relationships of Microsestean white-eyes (Zosteropidae)
RT based on mitochondrial sequence data."
RL Submitted (JUL-1999) to the EMBL/GenBank/DBJ databases.
DR EMBL: AF168438; AAG1314.1; -.
DR InterPro: IPR002429; Cyt_c_ox_2.
DR Pfam: PF00116; COX2; 1.
KW Mitochondrion.
FT NON_TER
FT SEQUENCE 29 AA; 3198 MW; 43F35FDA214CBD79 CRC64;

Query Match
Best Local Similarity 50.0%; Score 26; DB 8; Length 29;
Matches 5; Conservative 1; Mismatches 4; Indels 0; Gaps 0;

QY 1 PVVAESPCKP 10
DB 7 PIVVESAPLP 16

RESULT 11
084028 PRELIMINARY; PRT; 47 AA.
AC 084028;
DT 01-NOV-1996 (TREMBLrel. 01, Created)
DT 01-NOV-1996 (TREMBLrel. 01, Last sequence update)
DT 01-DEC-2001 (TREMBLrel. 19, Last annotation update)
DE INFLUENZA A/PUERTO RICO/8/34 (H0N1), MATRIX PROTEIN (SEG 7), RNA, 5'
DE END (FRAGMENT).
OS Influenza A virus.
OC Viruses; ssRNA negative-strand viruses; Orthomyxoviridae;
OC Influenza virus A and B group; Influenza A viruses.
OX NCBI_TaxID=11320;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=79213454; PubMed=572297;

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RA Both G.W., Air G.M.;
RT "Nucleotide sequence coding for the N-terminal region of the matrix
RT protein of influenza virus."
RL Eur. J. Biochem. 96:363-372(1979).
DR EMBL: M10642; AAA43309.1; -.
DR HSSP: P03485; IAA7.
DR InterPro: IPR001561; Flu_M1.
DR Pfam: PF00598; Flu_M1; 1.
DR Prodom: PD001061; Flu_M1; 1.
KW Matrix protein.
FT NON_TER
FT SEQUENCE 47 AA; 5343 MW; 83415B57C9CE4D51 CRC64;

Query Match
Best Local Similarity 50.0%; Score 26; DB 12; Length 47;
Matches 4; Conservative 2; Mismatches 3; Indels 0; Gaps 0;

QY 2 VVAESPCKP 10
DB 14 IVPSAPSKP 22

RESULT 12
096L04 PRELIMINARY; PRT; 18 AA.
AC 096L04;
DT 01-DEC-2001 (TREMBLrel. 19, Created)
DT 01-DEC-2001 (TREMBLrel. 19, Last sequence update)
DE 01-DEC-2001 (TREMBLrel. 19, Last annotation update)
DE HYPOTHETICAL 2.1 KDA PROTEIN (FRAGMENT).
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RA Strussberg R.;
RT Submitted (SEP-2001) to the EMBL/GenBank/DBJ databases.
DR EMBL: BC014606; AAHL4606.1; -.
KW Hypothetical protein.
FT NON_TER
FT SEQUENCE 18 AA; 2097 MW; 5165856AB840D268 CRC64;

Query Match
Best Local Similarity 48.1%; Score 25; DB 4; Length 18;
Matches 4; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

QY 4 AESPKP 10
DB 12 AESPRSP 18

RESULT 13
Q16070 PRELIMINARY; PRT; 20 AA.
AC Q16070;
DT 01-NOV-1996 (TREMBLrel. 01, Created)
DT 01-NOV-1996 (TREMBLrel. 01, Last sequence update)
DT 01-NOV-1998 (TREMBLrel. 08, Last annotation update)
DE NEUROFILAMENT HEAVY SUBUNIT (FRAGMENT).
GN NEFH.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=94040777; PubMed=8224877;
RA Figlewicz D.A., Rouleau G.A., Krizus A., Julien J.P.;
RT "Polymorphism in the multi-phosphorylation domain of the human

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RT neurofilament heavy-subunit-encoding gene.;
RL Gene 132:297-300(1983).
DR EMBL; S66488; AAB28609.1; -
FT NON_TER 1 1
SQ SEQUENCE 20 AA; 2198 MW; E9A0975B41FD8082 CRC64;

Query Match 48.1%; Score 25; DB 4; Length 20;
Best Local Similarity 55.6%; Pred. No. 7.2e+02;
Matches 5; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

OY 1 PVVASEPK 9
1 1 1 1 1 1
DB 2 PEKAKSPEK 10

RESULT 14

O9SLV6 PRELIMINARY; PRT; 22 AA.
ID O9SLV6
AC O9SLV6;
DT 01-MAY-2000 (TReMBLrel. 13, Created)
DT 01-MAY-2000 (TReMBLrel. 13, last sequence update)
DT 01-DEC-2001 (TReMBLrel. 19, last annotation update)
DE NITRILASE-LIKE PROTEIN (FRAGMENT).
GN TNIT4B.
OS Nicotiana tabacum (Common tobacco).
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots;
OC Asteridae; euasterids I; Solanales; Solanaceae; Nicotiana.
OX NCBI_TaxID=4097;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=SR-1;
RA MEDLINE=20039615; Pubmed=10574458;
RX Dohmoto M., Sano J., Tsunoda H., Yamaguchi K.;
RT "Structural analysis of the TNIT4 genes encoding nitrilase-like
RT protein from tobacco."
RL DNA Res. 6:313-317(1999).
DR EMBL; AB027127; BAA7683.1; -
FT NON_TER 22
SQ SEQUENCE 22 AA; 2320 MW; ACB9F59BC2461022 CRC64;

Query Match 48.1%; Score 25; DB 10; Length 22;
Best Local Similarity 71.4%; Pred. No. 7.9e+02;
Matches 5; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

OY 1 PVVASEP 7
1 1 1 1 1
DB 7 PVVNEGP 13

RESULT 15

O9UMH7 PRELIMINARY; PRT; 27 AA.
ID O9UMH7
AC O9UMH7;
DT 01-MAY-2000 (TReMBLrel. 13, Created)
DT 01-MAY-2000 (TReMBLrel. 13, last sequence update)
DT 01-DEC-2001 (TReMBLrel. 19, last annotation update)
DE SS-B/LA PROTEIN (FRAGMENT).
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=90237237; Pubmed=1692037;
RA Kohsaka H., Yamamoto K., Fujii H., Miura H., Miyasaka N., Nishioaka K.,
RA Miyamoto T.;
RT "Fine epitope mapping the human SS-B/La protein: Identification of a
RT distinct autoepitope homologous to a viral gag polypeptide."
RL J. Clin. Invest. 85:1566-1574(1990).
DR EMBL; M35261; AAA3652.1; -

FT NON_TER 1 1
FT NON_TER 27 27
SQ SEQUENCE 27 AA; 3175 MW; A9B8FED9F097F035 CRC64;

Query Match 48.1%; Score 25; DB 4; Length 27;
Best Local Similarity 50.0%; Pred. No. 9.7e+02;
Matches 4; Conservative 1; Mismatches 3; Indels 0; Gaps 0;

OY 3 VASEPKP 10
1 1 1 1 1 1
DB 9 IRRSPSKP 16

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Job time: 187 sec

GenCore version 4.5
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OM protein - protein search, using sw model

Run on: October 12, 2002, 16:51:25 ; Search time 60.75 Seconds

(without alignments)
18.284 Million cell updates/sec

Title: US-09-408-578a-1

Perfect score: 52

Sequence: 1 PVAESPCKP 10

Scoring table:

BLOSUM62
Gapop 10.0, Gapext 0.5

Searched: 747574 seqs, 11073796 residues

Total number of hits satisfying chosen parameters: 747574

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Database: A.GeneSeq_032802.*

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22: /SIDSL/gcgdata/geneSeq/geneSeq-emb1/AA2001.DAT.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	52	100.0	10	AAW41844	Modified B. burgdo
2	52	100.0	10	AAW41821	B. burgdorferi sen
3	52	100.0	11	AAW41825	Modified B. burgdo
4	52	100.0	15	AAW41827	B. burgdorferi sen
5	52	100.0	20	AAW41826	B. burgdorferi sen
6	52	100.0	23	AAW27428	B. burgdorferi sen
7	52	100.0	24	AAW27429	B. burgdorferi sen
8	52	100.0	19	AAW60897	Borrelia outer sur
9	52	100.0	19	AAW60897	Borrelia outer sur
10	52	100.0	19	AAW60897	Borrelia outer sur
11	52	100.0	19	AAW60897	Borrelia outer sur

12	52	100.0	194	15	AAW60894	Borrelia H9 antigen
13	52	100.0	194	15	AAW60896	Borrelia JSB antigen
14	52	100.0	194	21	AAW78908	Outer surface prot
15	52	100.0	206	22	AAW62723	B. burgdorferi str
16	52	100.0	207	16	AAW75730	B. burgdorferi str
17	52	100.0	207	18	AAW41823	B. burgdorferi str
18	52	100.0	209	16	AAW75728	B. burgdorferi str
19	52	100.0	209	22	AAW62730	B. burgdorferi str
20	52	100.0	210	16	AAW19335	B. burgdorferi str
21	52	100.0	210	16	AAW75727	Outer surface prot
22	52	100.0	211	22	AAW62722	B. burgdorferi str
23	52	100.0	212	12	AAW13140	B. burgdorferi str
24	52	100.0	212	16	AAW75729	B. burgdorferi str
25	52	100.0	212	18	AAW41824	B. burgdorferi str
26	52	100.0	377	22	AAW62713	B. burgdorferi str
27	52	100.0	378	22	AAW62712	B. burgdorferi str
28	52	100.0	378	22	AAW62725	B. burgdorferi str
29	52	100.0	384	22	AAW62726	B. burgdorferi str
30	52	100.0	386	22	AAW62727	B. burgdorferi str
31	52	100.0	400	22	AAW62739	B. burgdorferi str
32	52	100.0	401	22	AAW62733	B. burgdorferi str
33	52	100.0	401	22	AAW62738	B. burgdorferi str
34	52	100.0	408	22	AAW62737	B. burgdorferi str
35	52	100.0	410	22	AAW62740	B. burgdorferi str
36	52	100.0	466	16	AAW75739	B. burgdorferi str
37	52	100.0	466	16	AAW75740	B. burgdorferi str
38	52	100.0	587	16	AAW75746	B. burgdorferi str
39	51	98.1	191	15	AAW60884	B. burgdorferi str
40	49	94.2	10	18	AAW41838	B. burgdorferi str
41	49	94.2	11	15	AAW70367	B. burgdorferi str
42	49	94.2	15	15	AAW70362	B. burgdorferi str
43	49	94.2	189	15	AAW60907	B. burgdorferi str
44	49	94.2	189	15	AAW60909	B. burgdorferi str
45	49	94.2	191	15	AAW60898	B. burgdorferi str

ALIGNMENTS

RESULT 1

AAW41844 standard; peptide; 10 AA.

AAW41844;

14-MAY-1998 (first entry)

Modified B. burgdorferi sensu lato OspC C-terminal peptide.

Sensu lato: outer surface protein C; OspC; diagnosis; Lyme disease; vaccine; infection.

Borrelia burgdorferi.

Synthetic.

Key Location/Qualifiers

Modified-site /label=amlated

WO9742221-A1.

13-NOV-1997.

02-MAY-1997; 97WO-DK00203.

02-MAY-1996; 96DK-0000526.

(STAT-3) SPATENS SERUMINSTITUT.

Holm A, Mathiesen MJ, Ostergaard S, Theisen M;

WPI; 1997-558908/51.

PT Detecting previous sensitisation to the OspC protein of Borrelia
 PT burgdorferi - by detecting immunoreactivity between patient T cells
 or immunoglobulins and C-terminal peptide of the protein

PS Example 3; Page 53; 95pp; English.

CC The present sequence was used in the development of a novel method
 CC for the identification of a patient's previous sensitisation to
 CC Borrelia burgdorferi sensu lato outer surface protein C (OspC).
 CC The method comprises reacting immunoglobulin (Ig) or T cells from
 CC the patient with a polypeptide of at most 60 amino acids containing
 CC a peptide with at least 50% identity to the B. burgdorferi derived
 CC sequence AAW41821, or its subsequences of at least 5 amino acids. The
 CC degree of immunological reactivity between the polypeptide and Ig
 CC or T cells is measured and significant reactivity is indicative of
 CC sensitisation.
 CC The method can be used to diagnose Lyme disease and is based on
 CC reactivity with antibodies against the OspC protein. The test can
 CC be done in vitro or in vivo, e.g. as a skin test. Vaccine
 CC compositions comprising the polypeptide can be used to protect
 CC humans and other animals against B. burgdorferi infection. The
 CC polypeptide provides higher sensitivity than full-length OspC, and
 CC so is better at detecting infection in its early stages, especially
 CC when combined with the known assay for flagellar proteins. The
 CC seven carboxy-terminal residues of AAW41821 represent an epitope
 CC essential for human immune response to OspC. The polypeptide is
 CC also easier to prepare and purify than (nearly) full-length
 CC protein, facilitating standardisation of the assay, and is less
 CC cross-reactive with antibodies raised against other antigens. The
 CC small size of the polypeptide allows a high density of binding
 CC sites to be created on a solid support. Incorporation of
 CC non-natural amino acid into the polypeptide increases its
 CC resistance to peptidases when used in vivo.

CC Sequence 10 AA;

Query Match 100.0%; Score 52; DB 18; Length 10;

Best Local Similarity 100.0%; Pred. No. 0.0078;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 PVVAESPCKP 10

DB 1 PVVAESPCKP 10

RESULT 2

ID AAW41821 standard; peptide; 10 AA.

AC AAW41821;

DT 14-MAY-1998 (first entry)

DE B. burgdorferi sensu lato OspC carboxy-terminal peptide.

KW Senu lato; outer surface protein C; OspC; diagnosis; Lyme disease;

KW vaccine; infection.

OS Borrelia burgdorferi.

PN WO9742221-A1.

PD 13-NOV-1997.

PF 02-MAY-1997; 97WO-DK00203.

PR 02-MAY-1996; 96DK-0000526.

PA (STAT-) STATENS SERUMINSTITUT.

XX Holm A, Mathiesen MJ, Ostergaard S, Thølsen M;

DR WPI: 1997-558908/51.

XX Detecting previous sensitisation to the OspC protein of Borrelia
 PT burgdorferi - by detecting immunoreactivity between patient T cells
 or immunoglobulins and C-terminal peptide of the protein

PS Claim 1; Page 77; 95pp; English.

CC The present sequence was used in the development of a novel method
 CC for the identification of a patient's previous sensitisation to
 CC Borrelia burgdorferi sensu lato outer surface protein C (OspC).
 CC The method comprises reacting immunoglobulin (Ig) or T cells from
 CC the patient with a polypeptide of at most 60 amino acids containing
 CC a peptide with at least 50% identity to the B. burgdorferi derived
 CC sequence AAW41821, or its subsequences of at least 5 amino acids. The
 CC degree of immunological reactivity between the polypeptide and Ig
 CC or T cells is measured and significant reactivity is indicative of
 CC sensitisation.
 CC The method can be used to diagnose Lyme disease and is based on
 CC reactivity with antibodies against the OspC protein. The test can
 CC be done in vitro or in vivo, e.g. as a skin test. Vaccine
 CC compositions comprising the polypeptide can be used to protect
 CC humans and other animals against B. burgdorferi infection. The
 CC polypeptide provides higher sensitivity than full-length OspC, and
 CC so is better at detecting infection in its early stages, especially
 CC when combined with the known assay for flagellar proteins. The
 CC seven carboxy-terminal residues of AAW41821 represent an epitope
 CC essential for human immune response to OspC. The polypeptide is
 CC also easier to prepare and purify than (nearly) full-length
 CC protein, facilitating standardisation of the assay, and is less
 CC cross-reactive with antibodies raised against other antigens. The
 CC small size of the polypeptide allows a high density of binding
 CC sites to be created on a solid support. Incorporation of
 CC non-natural amino acid into the polypeptide increases its
 CC resistance to peptidases when used in vivo.

CC Sequence 10 AA;

Query Match 100.0%; Score 52; DB 18; Length 10;

Best Local Similarity 100.0%; Pred. No. 0.0078;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 PVVAESPCKP 10

DB 1 PVVAESPCKP 10

RESULT 3

ID AAW41825 standard; peptide; 11 AA.

AC AAW41825;

DT 14-MAY-1998 (first entry)

DE Modified B. burgdorferi sensu lato OspC C-terminal peptide.

KW Senu lato; outer surface protein C; OspC; diagnosis; Lyme disease;

KW vaccine; infection.

OS Borrelia burgdorferi.

OS Synthetic.

FT Key Location/Qualifiers

FT Modified-site 1 /note- "6-aminohecanoic acid"

PN WO9742221-A1.

PD 13-NOV-1997.

PF 02-MAY-1997; 97WO-DK00203.

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XX 02-MAY-1996; 96DK-0000526.
PR (STAT-) STATENS SERUMINSTITUT.
PA Holm A, Mathiesen MJ, Ostergaard S, Thelsen M;
PI WPI: 1997-558908/51.
XX
XX Detecting previous sensitisation to the Ospc protein of Borellia
XX burgdorferi - by detecting immunoreactivity between patient T cells
XX or immunoglobulins and C-terminal peptide of the protein
XX
XX Example; Page 40; 95pp; English.
XX
XX The present sequence was used in the development of a novel method
XX for the identification of a patient's previous sensitisation to
XX Borellia burgdorferi sensu lato outer surface protein C (OspC).
XX The method comprises reacting immunoglobulin (Ig) or T cells from
XX the patient with a polypeptide of at most 60 amino acids containing
XX a peptide with at least 50% identity to the B. burgdorferi derived
XX sequence AAW41821, or its subsequences of at least 5 amino acids. The
XX degree of immunological reactivity between the polypeptide and Ig
XX or T cells is measured and significant reactivity is indicative of
XX sensitisation.
XX The method can be used to diagnose Lyme disease and is based on
XX reactivity with antibodies against the Ospc protein. The test can
XX be done in vitro or in vivo, e.g. as a skin test. Vaccine
XX compositions comprising the polypeptide can be used to protect
XX humans and other animals against B. burgdorferi infection. The
XX polypeptide provides higher sensitivity than full-length Ospc, and
XX so is better at detecting infection in its early stages, especially
XX when combined with the known assay for flagellar proteins. The
XX seven carboxy-terminal residues of AAW41821 represent an epitope
XX essential for human immune response to Ospc. The polypeptide is
XX also easier to prepare and purify than (nearly) full-length
XX protein, facilitating standardisation of the assay, and is less
XX cross-reactive with antibodies raised against other antigens. The
XX small size of the polypeptide allows a high density of binding
XX sites to be created on a solid support. Incorporation of
XX non-natural amino acid into the polypeptide increases its
XX resistance to peptidases when used in vivo.
XX
XX Sequence 11 AA:
SQ

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Query Match 100.0%; Score 52; DB 18; Length 11;
Best Local Similarity 100.0%; Pred. No. 0.0085;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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QY 1 PVVAESPCKP 10
   |||||||||
DB 2 PVVAESPCKP 11

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RESULT 4
AAW41827
ID AAW41827 standard; peptide; 15 AA.
XX
XX AAW41827;
XX
XX 14-MAY-1998 (first entry)
XX
XX B. burgdorferi sensu lato Ospc carboxy-terminal peptide.
XX
XX Sensu lato; outer surface protein C; Ospc; diagnosis; Lyme disease;
XX vaccine; infection.
XX
XX Borellia burgdorferi.
XX
XX WO9742221-A1.
XX
XX 13-NOV-1997.
PD

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XX 02-MAY-1997; 97WO-DK00203.
PF
XX
XX 02-MAY-1996; 96DK-0000526.
PR
XX
XX (STAT-) STATENS SERUMINSTITUT.
PA Holm A, Mathiesen MJ, Ostergaard S, Thelsen M;
PI WPI: 1997-558908/51.
XX
XX Detecting previous sensitisation to the Ospc protein of Borellia
XX burgdorferi - by detecting immunoreactivity between patient T cells
XX or immunoglobulins and C-terminal peptide of the protein
XX
XX Example 3; Page 51; 95pp; English.
XX
XX The present sequence was used in the development of a novel method
XX for the identification of a patient's previous sensitisation to
XX Borellia burgdorferi sensu lato outer surface protein C (OspC).
XX The method comprises reacting immunoglobulin (Ig) or T cells from
XX the patient with a polypeptide of at most 60 amino acids containing
XX a peptide with at least 50% identity to the B. burgdorferi derived
XX sequence AAW41821, or its subsequences of at least 5 amino acids. The
XX degree of immunological reactivity between the polypeptide and Ig
XX or T cells is measured and significant reactivity is indicative of
XX sensitisation.
XX The method can be used to diagnose Lyme disease and is based on
XX reactivity with antibodies against the Ospc protein. The test can
XX be done in vitro or in vivo, e.g. as a skin test. Vaccine
XX compositions comprising the polypeptide can be used to protect
XX humans and other animals against B. burgdorferi infection. The
XX polypeptide provides higher sensitivity than full-length Ospc, and
XX so is better at detecting infection in its early stages, especially
XX when combined with the known assay for flagellar proteins. The
XX seven carboxy-terminal residues of AAW41821 represent an epitope
XX essential for human immune response to Ospc. The polypeptide is
XX also easier to prepare and purify than (nearly) full-length
XX protein, facilitating standardisation of the assay, and is less
XX cross-reactive with antibodies raised against other antigens. The
XX small size of the polypeptide allows a high density of binding
XX sites to be created on a solid support. Incorporation of
XX non-natural amino acid into the polypeptide increases its
XX resistance to peptidases when used in vivo.
XX
XX Sequence 15 AA:
SQ

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```

Query Match 100.0%; Score 52; DB 18; Length 15;
Best Local Similarity 100.0%; Pred. No. 0.012;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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QY 1 PVVAESPCKP 10
   |||||||||
DB 6 PVVAESPCKP 15

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```

RESULT 5
AAW41826
ID AAW41826 standard; peptide; 20 AA.
XX
XX AAW41826;
XX
XX 14-MAY-1998 (first entry)
XX
XX B. burgdorferi sensu lato Ospc carboxy-terminal peptide.
XX
XX Sensu lato; outer surface protein C; Ospc; diagnosis; Lyme disease;
XX vaccine; infection.
XX
XX Borellia burgdorferi.
XX
XX WO9742221-A1.
XX
XX
XX

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XX 13-NOV-1997.
 PD
 XX 02-MAY-1997; 97WO-DK00203.
 PF
 XX 02-MAY-1996; 96DK-0000526.
 PR
 XX (STAT-) STATENS SERUMINSTITUT.
 PA
 XX Holm A, Mathiesen MJ, Ostergaard S, Theisen M;
 PI WPI; 1997-558908/51.
 DR
 XX
 PT Detecting previous sensitisation to the OspC protein of *Borrelia*
 PT burgdorferi - by detecting immunoreactivity between patient T cells
 PT or immunoglobulins and C-terminal peptide of the protein
 PS
 XX
 XX Example 3; Page 51; 95pp; English.
 CC The present sequence was used in the development of a novel method
 CC for the identification of a patient's previous sensitisation to
 CC *Borrelia burgdorferi sensu lato* outer surface protein C (OspC).
 CC The method comprises reacting immunoglobulin (Ig) or T cells from
 CC the patient with a polypeptide of at most 60 amino acids containing
 CC a peptide with at least 50% identity to the B. burgdorferi derived
 CC sequence AAW41821, or its subsequences of at least 5 amino acids. The
 CC degree of immunological reactivity between the polypeptide and Ig
 CC or T cells is measured and significant reactivity is indicative of
 CC sensitisation.
 CC The method can be used to diagnose Lyme disease and is based on
 CC reactivity with antibodies against the OspC protein. The test can
 CC be done *in vitro* or *in vivo*, e.g. as a skin test. Vaccine
 CC compositions comprising the polypeptide can be used to protect
 CC humans and other animals against B. burgdorferi infection. The
 CC polypeptide provides higher sensitivity than full-length OspC, and
 CC so is better at detecting infection in its early stages, especially
 CC when combined with the known assay for flagellar proteins. The
 CC seven carboxy-terminal residues of AAW41821 represent an epitope
 CC essential for human immune response to OspC. The polypeptide is
 CC also easier to prepare and purify than (nearly) full-length
 CC protein, facilitating standardisation of the assay, and is less
 CC cross-reactive with antibodies raised against other antigens. The
 CC small size of the polypeptide allows a high density of binding
 CC sites to be created on a solid support. Incorporation of
 CC non-natural amino acid into the polypeptide increases its
 CC resistance to peptidases when used *in vivo*.
 XX
 SQ Sequence 20 AA:
 Query Match 100.0%; Score 52; DB 18; Length 20;
 Best Local Similarity 100.0%; Pred. No. 0.016;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0.
 QY 1 PVVAESPCKP 10
 |||||||||
 DB 11 PVVAESPCKP 20
 RESULT 6
 AAY27428
 ID AAY27428 standard; peptide: 23 AA.
 XX
 XX AAY27428;
 XX
 XX 26-NOV-1999 (first entry)
 XX
 XX *Borrelia* outer surface protein C (OspC) C-terminal peptide fragment.
 XX
 XX *Borrelia*; Igm antibody; outer surface protein C; Ospc; deer tick;
 KW cerebrospinal fluid; Lyme borreliosis; micro-capture assay; flagellum;
 epitope.
 XX

OS	Synthetic.
XS	Borrelia burgdorferi.
XK	
XX	Key Location/Qualifiers
FH	Modified-site 1 /note= "linked to biotin via an O-linker of formula [2-aminoethoxy]ethoxy acetic acid"
FT	
ET	
PV	
NN	EP949508-A1.
PD	13-OCT-1999.
XP	
PE	07-APR-1999; 99EP-0610026.
PR	08-APR-1998; 98DK-0000516.
PA	(DAKO-) DAKO AS.
PI	
PL	Stafieldt Schou O., Winther L., Stender H;
DR	WPI; 1999-553537/47.
PT	
PF	Diagnosing Lyme borreliosis by detecting antibodies against two antigens simultaneously -
PS	
PP	Example I; Page 7; 23pp; English.
CC	The invention provides a new method for detecting IgM antibodies against Borrelia burgdorferi in a sample of human or animal fluid. The method comprises: (1) contacting antibodies in the sample with anti-IgM immobilized to a solid support, (2) separating the support from the liquid phase; and (3) contacting the bound antibodies with a complex comprising at least one set of B. burgdorferi outer surface protein C (OsPC) peptides and/or at least one set of other B. burgdorferi peptides, each attached to a carrier; and (4) detecting the presence of antibodies against B. burgdorferi. The new method may be used to detect antibodies against B. burgdorferi in (especially) serum or cerebrospinal fluid samples from patients bitten by deer ticks. B. burgdorferi causes Lyme borreliosis so detection of antibodies against it allows diagnosis of infection by this organism. The method is a micro-capture assay in which the antigen complex is a combination of the B. burgdorferi flagellum and OsPC peptides. The presence of epitopes from both antigens in the complex allows the simultaneous detection of serum antibodies against these proteins which increases the sensitivity of the test. The two antigens are pure, which also decreases the possibility of cross-reactivity. The present sequence represents a Borrelia OspC C-terminal peptide fragment.
CC	
CC	
CX	Sequence 23 AA:
SQ	
Query Match	100.0%; Score 52; DB 20; Length 23;
Best Local Similarity	100.0%; Prid. NO. 0.018;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;	
OY	1 PVVAESPCKP 10
Dd	14 pvaespckp 23
RESULT 7	
AAY27429	
ID AAY27429 standard; peptide: 24 AA.	
AAI27429;	
DT 26-NOV-1999 (first entry)	
DE	Borrelia outer surface protein c (ospc) C-terminal peptide fragment.
KW	Borrelia; IgM antibody; outer surface protein C; OsPC; deer tick; cerebrospinal fluid; Lyme borreliosis; micro-capture assay; flagellum; epitope.

OS Synthetic.
 OS Borrelia burgdorferi.
 XX
 PN EP949508-A1.
 XX
 PD 13-OCT-1999.
 XX
 PF 07-APR-1999; 99EP-0610026.
 XX
 PR 08-APR-1998; 98DK-0000516.
 XX
 PA (DAKO-) DAKO AS.
 PI Staffeldt Schou O, Winther L, Stender H;
 XX
 DR WPI; 1999-553537/47.
 XX
 PT Diagnosing Lyme borreliosis by detecting antibodies against two
 PT antigens simultaneously -
 XX
 PS Example 1; Page 7; 23pp; English.
 XX
 CC The invention provides a new method for detecting IgM antibodies against
 CC Borrelia burgdorferi in a sample of human or animal fluid. The method
 CC comprises: (1) contacting antibodies in the sample with anti-IgM
 CC immobilized to a solid support, (2) separating the support from the
 CC liquid phase; and (3) contacting the bound antibodies with a complex
 CC comprising at least one set of B. burgdorferi (outer surface protein C
 CC (OspC) peptides and/or at least one set of other B. burgdorferi peptides,
 CC each attached to a carrier; and (4) detecting the presence of antibodies
 CC against B. burgdorferi. The new method may be used to detect antibodies
 CC against B. burgdorferi in (especially) serum or cerebrospinal fluid
 CC samples from patients bitten by deer ticks. B. burgdorferi causes Lyme
 CC borreliosis so detection of antibodies against it allows diagnosis of
 CC infection by this organism. The method is a micro-capture assay in which
 CC the antigen complex is a combination of the B. burgdorferi flagellum and
 CC OspC peptides. The presence of epitopes from both antigens in the complex
 CC allows the simultaneous detection of serum antibodies against these
 CC proteins which increases the sensitivity of the test. The two antigens
 CC are pure, which also decreases the possibility of cross-reactivity. The
 CC present sequence represents a Borrelia OspC C-terminal peptide fragment,
 CC where the N-terminal cysteine residue has been incorporated to provide a
 CC SH group to be used in a coupling reaction.
 CC
 SQ Sequence 24 AA;
 XX
 XX
 Query Match 100.0%; Score 52; DB 20; Length 24;
 Best Local Similarity 100.0%; Pred. No. 0.019;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 PVVAESPCKP 10
 |||||||||
 DB 15 PVVAESPCKP 24
 RESULT 8
 AAR60897
 ID AAR60897 standard; Protein; 191 AA.
 XX
 AC AAR60897;
 XX
 DT 25-MAY-1995 (first entry)
 XX
 DE Borrelia VS461 antigen vaccine.
 XX
 KW OSpC antigen; vaccine; Lyme disease; borreliosis; immunogen;
 KW serovar typing; restriction fragment length polymorphism;
 KW RFLP; Pichia pastoris.
 XX
 OS Borrelia burgdorferi VS461.
 OS
 PN WO9425596-A.
 XX

XX
 PD 10-NOV-1994.
 XX
 PF 29-APR-1994; 94WO-EP01365.
 XX
 PR 29-APR-1993; 93US-0053863.
 XX
 PA (IMMO) IMMUNO AG.
 XX
 PI Crowe B, Dornier F, Lavey I;
 XX
 DR WPI; 1994-358273/44.
 DR N-PSDB; AAQ73870.
 XX
 PT Immunogenic composition comprising OSpC antigens - for the
 PT treatment of Lyme borreliosis in different, specific geographical
 PT areas.
 XX
 PS Disclosure; Fig. 9a; 115pp; English.
 XX
 CC A vaccine for Lyme disease includes selected OSpC antigen
 CC formulations based on defined OSpC families resolved by serovar
 CC typing and RFLP typing. Partial sequences of OSpC genes selected
 CC from different RFLP types are given in AAQ73883-905 (encoded peptides,
 CC comprising the first 92% of mature OSpC, are given in AAR62771-93).
 CC Complete sequences of these novel OSpC genes, including the 3' end,
 CC plus sequences for the OSpC genes of Borrelia strains H13 and 28691
 CC are given in AAQ73857-82, and encoded proteins in AAR60884-909. The
 CC DNA sequences may be expressed in e.g. Pichia pastoris for
 CC recombinant antigen production.
 CC
 SQ Sequence 191 AA;
 XX
 XX
 Query Match 100.0%; Score 52; DB 15; Length 191;
 Best Local Similarity 100.0%; Pred. No. 0.15;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 PVVAESPCKP 10
 |||||||||
 DB 182 PVVAESPCKP 191
 RESULT 9
 AAR60889
 ID AAR60889 standard; Protein; 192 AA.
 XX
 AC AAR60889;
 XX
 DT 25-MAY-1995 (first entry)
 XX
 DE Borrelia 297 antigen vaccine.
 XX
 KW OSpC antigen; vaccine; Lyme disease; borreliosis; immunogen;
 KW serovar typing; restriction fragment length polymorphism;
 KW RFLP; Pichia pastoris.
 XX
 OS Borrelia burgdorferi 297.
 OS
 PN WO9425596-A.
 XX
 PD 10-NOV-1994.
 XX
 PF 29-APR-1994; 94WO-EP01365.
 XX
 PR 29-APR-1993; 93US-0053863.
 XX
 PA (IMMO) IMMUNO AG.
 XX
 PI Crowe B, Dornier F, Lavey I;
 XX
 DR WPI; 1994-358273/44.
 DR N-PSDB; AAQ73862.
 XX

XX Immunogenic composition comprising OspC antigens - for the
 PT treatment of Lyme borreliosis in different, specific geographical
 PT areas.
 PS
 XX
 PS Disclosure; Fig. 9a; 115pp; English.
 CC
 CC A vaccine for Lyme disease includes selected OspC antigen
 CC formulations based on defined OspC families resolved by serovar
 CC typing and RFLP typing. Partial sequences of OspC genes selected
 CC from different RFLP types are given in AAQ73883-905 (encoded peptides,
 CC comprising the first 92% of mature OspC, are given in AAR62771-93).
 CC Complete sequences of these novel ospc genes, including the 3' end,
 CC plus sequences for the ospc genes of Borrelia strains H13 and 28691
 CC are given in AAQ73857-82, and encoded proteins in AAR60884-909. The
 CC DNA sequences may be expressed in e.g. Pichia pastoris for
 CC recombinant antigen production.
 CC
 SQ Sequence 192 AA;
 Query Match 100.0%; Score 52; DB 15; Length 192;
 Best Local Similarity 100.0%; Pred. No. 0.15;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 PVTAESEPKP 10
 DB 183 pvtasepkkp 192
 RESULT 10
 AAR60886
 ID AAR60886 standard; Protein; 192 AA.
 AC AAR60886;
 XX
 XX
 DT 25-MAY-1995 (first entry)
 DE Borrelia IP2 OspC antigen vaccine.
 XX
 XX
 KM OSpC antigen; vaccine; Lyme disease; borreliosis; immunogen;
 KM serovar typing; restriction fragment length polymorphism;
 KM RFLP; Pichia pastoris.
 XX
 OS Borrelia burgdorferi IP2.
 OS
 XX WO9425596-A.
 PN
 XX 10-NOV-1994.
 PD
 XX
 PF 29-APR-1994; 94WO-EP01365.
 XX
 PR 29-APR-1993; 93US-0053863.
 XX
 PA (IMMO) IMMUNO AG.
 PA
 PI Crowe B, Dornier F, Livey I;
 PI
 XX WPI; 1994-358273/44.
 DR N-PSDB; AAQ73859.
 DR
 XX
 PT Immunogenic composition comprising OspC antigens - for the
 PT treatment of Lyme borreliosis in different, specific geographical
 PT areas.
 PS
 XX Disclosure; Fig. 9a; 115pp; English.
 PS
 CC A vaccine for Lyme disease includes selected OspC antigen
 CC formulations based on defined OspC families resolved by serovar
 CC typing and RFLP typing. Partial sequences of OspC genes selected
 CC from different RFLP types are given in AAQ73883-905 (encoded peptides,
 CC comprising the first 92% of mature OspC, are given in AAR62771-93).
 CC Complete sequences of these novel ospc genes, including the 3' end,
 CC plus sequences for the ospc genes of Borrelia strains H13 and 28691
 CC are given in AAR62771-93).
 CC Complete sequences of these novel ospc genes, including the 3' end,
 CC

CC plus sequences for the ospc genes of Borrelia strains H13 and 28691
 CC are given in AAQ73857-82, and encoded proteins in AAR60884-909. The
 CC DNA sequences may be expressed in e.g. Pichia pastoris for
 CC recombinant antigen production.
 CC
 SQ Sequence 192 AA;
 Query Match 100.0%; Score 52; DB 15; Length 192;
 Best Local Similarity 100.0%; Pred. No. 0.15;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 PVTAESEPKP 10
 DB 183 pvtasepkkp 192
 RESULT 11
 AAR60893
 ID AAR60893 standard; Protein; 193 AA.
 AC AAR60893;
 XX
 XX
 DT 25-MAY-1995 (first entry)
 DE Borrelia ACA1 antigen vaccine.
 XX
 XX
 KM OSpC antigen; vaccine; Lyme disease; borreliosis; immunogen;
 KM serovar typing; restriction fragment length polymorphism;
 KM RFLP; Pichia pastoris.
 XX
 OS Borrelia burgdorferi ACA1.
 OS
 XX WO9425596-A.
 PN
 XX 10-NOV-1994.
 PD
 XX
 PF 29-APR-1994; 94WO-EP01365.
 XX
 PR 29-APR-1993; 93US-0053863.
 XX
 PA (IMMO) IMMUNO AG.
 PA
 PI Crowe B, Dornier F, Livey I;
 PI
 XX WPI; 1994-358273/44.
 DR N-PSDB; AAQ73866.
 DR
 XX
 PT Immunogenic composition comprising OspC antigens - for the
 PT treatment of Lyme borreliosis in different, specific geographical
 PT areas.
 PS
 XX Disclosure; Fig. 9a; 115pp; English.
 PS
 CC A vaccine for Lyme disease includes selected OspC antigen
 CC formulations based on defined OspC families resolved by serovar
 CC typing and RFLP typing. Partial sequences of OspC genes selected
 CC from different RFLP types are given in AAQ73883-905 (encoded peptides,
 CC comprising the first 92% of mature OspC, are given in AAR62771-93).
 CC Complete sequences of these novel ospc genes, including the 3' end,
 CC plus sequences for the ospc genes of Borrelia strains H13 and 28691
 CC are given in AAQ73857-82, and encoded proteins in AAR60884-909. The
 CC DNA sequences may be expressed in e.g. Pichia pastoris for
 CC recombinant antigen production.
 CC
 SQ Sequence 193 AA;
 Query Match 100.0%; Score 52; DB 15; Length 193;
 Best Local Similarity 100.0%; Pred. No. 0.15;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 PVTAESEPKP 10

Db 184 pvaesppk 193

|||||

RESULT 12

AA60894 ID AAR60894 standard; Protein: 194 AA.

AC AAR60894;

DT 25-MAY-1995 (first entry)

DE Borrelia H9 antigen vaccine.

OSPC antigen; vaccine; Lyme disease; borreliosis; immunogen;

KW serovar typing; restriction fragment length polymorphism;

KM RFLP; Pichia pastoris.

XX Borrelia burgdorferi H9.

XX WO9425596-A.

XX 10-NOV-1994.

XX 29-APR-1994; 94WO-EP01365.

XX 29-APR-1993; 93US-0053863.

XX (IMMO) IMMUNO AG.

XX Crowe B, Dorner F, Lively I;

XX WPI; 1994-358273/44.

XX DR N-PSDB; AAQ60894.

XX Immunogenic composition comprising Ospc antigens - for the

XX treatment of Lyme borreliosis in different, specific geographical

XX areas.

XX Disclosure; Fig. 9a; 115pp; English.

XX A vaccine for Lyme disease includes selected Ospc antigen

XX formulations based on defined Ospc families resolved by serovar

XX typing and RFLP typing. Partial sequences of Ospc genes selected

XX from different RFLP types are given in AAQ73883-905 (encoded peptides,

XX comprising the first 92% of mature Ospc, are given in AAR62771-93).

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XX are given in AAQ73857-82, and encoded proteins in AAR60884-909. The

XX DNA sequences may be expressed in e.g. Pichia pastoris for

XX recombinant antigen production.

XX Sequence 194 AA;

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XX OSPC antigen; vaccine; Lyme disease; borreliosis; immunogen;

XX serovar typing; restriction fragment length polymorphism;

XX RFLP; Pichia pastoris.

XX Borrelia burgdorferi USB.

XX WO9425596-A.

XX 10-NOV-1994.

XX 29-APR-1994; 94WO-EP01365.

XX 29-APR-1993; 93US-0053863.

XX (IMMO) IMMUNO AG.

XX Crowe B, Dorner F, Lively I;

XX WPI; 1994-358273/44.

XX DR N-PSDB; AAQ73869.

XX Immunogenic composition comprising Ospc antigens - for the

XX treatment of Lyme borreliosis in different, specific geographical

XX areas.

XX Disclosure; Fig. 9a; 115pp; English.

XX A vaccine for Lyme disease includes selected Ospc antigen

XX formulations based on defined Ospc families resolved by serovar

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XX Sequence 194 AA;

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XX OSPC antigen; vaccine; Lyme disease; borreliosis; immunogen;

XX serovar typing; restriction fragment length polymorphism;

XX RFLP; Pichia pastoris.

XX Borrelia burgdorferi USB.

XX WO9425596-A.

XX 10-NOV-1994.

XX 29-APR-1994; 94WO-EP01365.

XX 29-APR-1993; 93US-0053863.

XX (IMMO) IMMUNO AG.

XX Crowe B, Dorner F, Lively I;

XX WPI; 1994-358273/44.

XX DR N-PSDB; AAQ73869.

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XX Disclosure; Fig. 9a; 115pp; English.

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XX Sequence 194 AA;

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XX OSPC antigen; vaccine; Lyme disease; borreliosis; immunogen;

XX serovar typing; restriction fragment length polymorphism;

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XX Borrelia burgdorferi USB.

XX WO9425596-A.

XX 10-NOV-1994.

XX 29-APR-1994; 94WO-EP01365.

XX 29-APR-1993; 93US-0053863.

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XX Crowe B, Dorner F, Lively I;

XX WPI; 1994-358273/44.

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XX Sequence 194 AA;

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XX OSPC antigen; vaccine; Lyme disease; borreliosis; immunogen;

XX serovar typing; restriction fragment length polymorphism;

XX RFLP; Pichia pastoris.

XX Borrelia burgdorferi USB.

XX WO9425596-A.

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XX 29-APR-1994; 94WO-EP01365.

XX 29-APR-1993; 93US-0053863.

XX (IMMO) IMMUNO AG.

XX Crowe B, Dorner F, Lively I;

XX WPI; 1994-358273/44.

XX DR N-PSDB; AAQ73869.

XX Immunogenic composition comprising Ospc antigens - for the

XX treatment of Lyme borreliosis in different, specific geographical

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XX Disclosure; Fig. 9a; 115pp; English.

XX A vaccine for Lyme disease includes selected Ospc antigen

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XX DNA sequences may be expressed in e.g. Pichia pastoris for

XX recombinant antigen production.

XX Sequence 194 AA;

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XX OSPC antigen; vaccine; Lyme disease; borreliosis; immunogen;

XX serovar typing; restriction fragment length polymorphism;

XX RFLP; Pichia pastoris.

XX Borrelia burgdorferi USB.

XX WO9425596-A.

XX 10-NOV-1994.

XX 29-APR-1994; 94WO-EP01365.

XX 29-APR-1993; 93US-0053863.

XX (IMMO) IMMUNO AG.

XX Crowe B, Dorner F, Lively I;

XX WPI; 1994-358273/44.

XX DR N-PSDB; AAQ73869.

XX Immunogenic composition comprising Ospc antigens - for the

XX treatment of Lyme borreliosis in different, specific geographical

XX areas.

XX Disclosure; Fig. 9a; 115pp; English.

XX A vaccine for Lyme disease includes selected Ospc antigen

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XX from different RFLP types are given in AAQ73883-905 (encoded peptides,

XX comprising the first 92% of mature Ospc, are given in AAR62771-93).

XX Complete sequences of these novel ospc genes, including the 3' end,

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XX are given in AAQ73857-82, and encoded proteins in AAR60884-909. The

XX DNA sequences may be expressed in e.g. Pichia pastoris for

XX recombinant antigen production.

XX Sequence 194 AA;

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XX OSPC antigen; vaccine; Lyme disease; borreliosis; immunogen;

XX serovar typing; restriction fragment length polymorphism;

XX RFLP; Pichia pastoris.

XX Borrelia burgdorferi USB.

XX WO9425596-A.

XX 10-NOV-1994.

XX 29-APR-1994; 94WO-EP01365.

XX 29-APR-1993; 93US-0053863.

XX (IMMO) IMMUNO AG.

XX Crowe B, Dorner F, Lively I;

XX WPI; 1994-358273/44.

XX DR N-PSDB; AAQ73869.

XX Immunogenic composition comprising Ospc antigens - for the

XX treatment of Lyme borreliosis in different, specific geographical

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XX Disclosure; Fig. 9a; 115pp; English.

XX A vaccine for Lyme disease includes selected Ospc antigen

XX formulations based on defined Ospc families resolved by serovar

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XX from different RFLP types are given in AAQ73883-905 (encoded peptides,

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XX are given in AAQ73857-82, and encoded proteins in AAR60884-909. The

XX DNA sequences may be expressed in e.g. Pichia pastoris for

XX recombinant antigen production.

XX Sequence 194 AA;

XX

XX

XX

XX (GUND-) GUNDERSEN LUTHERAN MEDICAL FOUND INC.
 XX Callister SN, Lovrich SD, Schell RF, Jobe DA;
 XX MPI; 2000-195305/17.
 DR N-PSDB; AAF292216.
 XX
 PT New immunogenic polypeptides useful as a vaccine against Lyme disease
 PT and for treating and detecting Borrelia infection in mammals consists
 PT an epitope of Borrelia burgdorferi OspC fragment.
 PS
 XX Claim 3: Fig 4; 51pp; English.
 CC This sequence represents the Borrelia burgdorferi outer surface protein C
 CC (OspC) DraI fragment amino acid sequence. The polypeptide contains an
 CC immunological epitope used in the invention. Large amounts of OspC are
 CC rapidly synthesised by B. burgdorferi shortly after attachment of
 CC infected ticks to mammalian hosts. The OspC protein sequence is used to
 CC diagnose B. borrelia infection in mammals. The OspC nucleotide sequence
 CC is used to prevent (via vaccination), treat or detect Borrelia
 CC (especially B. burgdorferi) infections, i.e. Lyme disease, in mammals
 CC including humans. The OspC nucleotide sequence provides a superior
 CC diagnostic antigen that detects early Lyme disease infection, predicts
 CC successful eradication or the organism from the host, and discriminates
 CC between individuals with Lyme disease and individuals who have been
 CC vaccinated with an OspA Lyme disease vaccination. Detection of anti-OspC
 CC borrelial antibodies advantageously gives an early diagnosis which
 CC anti-OspA and anti-OspB borrelial antibodies cannot do. Unlike
 CC vaccination with OspA, vaccination with OspC results in clearance of
 CC spirochetes and resolution of symptoms even if administered after
 CC infection with B. burgdorferi.
 XX
 SQ Sequence 194 AA;

Query Match 100.0%; Score 52; DB 21; Length 194;
 Best Local Similarity 100.0%; Pred. No. 0.15;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 PVTAESEPKKP 10
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 DB 185 pvtasepkkp 194

RESULT 15
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 ID AAB62723 standard; Protein; 206 AA.
 XX
 AC AAB62723;
 XX
 DT 03-APR-2001 (first entry)
 XX
 DE B afzeili ospc protein SEQ ID NO: 50.
 XX
 KM Borrelia; ospc; Lyme disease; vaccine; chimeric protein; tick.
 XX
 OS Borrelia afzeili.
 XX
 PN WO200078966-A1.
 XX
 PD 28-DEC-2000.
 XX
 PF 19-JUN-2000; 2000WO-US16915.
 XX
 PR 18-JUN-1999; 99US-0140042.
 XX
 PA (UANY) UNIV NEW YORK STATE RES FOUND.
 PA (BROO-) BROOK BIOTECHNOLOGIES INC.
 XX
 PI Dattwyler RJ, Seino G, Dykhuizen D, Luft BJ, Gomes-Solecki M;
 XX
 DR MPI; 2001-050113/06.

DR N-PSDB; AAF29027.
 XX
 PT Compositions of OspC polypeptides from strains of Borrelia which cause
 PT Lyme disease are used to immunize animals and detect immune responses
 PT to Lyme disease.
 PS
 XX Disclosure; Page 109; 160pp; English.

CC The present invention provides compositions comprising ospc proteins and
 CC chimeric ospc proteins from members of the Borrelia genus. These may be
 CC Borrelia burgdorferi, B. afzeili or B. garinii. These can be used as
 CC vaccines against Borrelia infection, which is spread by ticks and leads
 CC to Lyme disease.
 XX
 SQ Sequence 206 AA;

Query Match 100.0%; Score 52; DB 22; Length 206;
 Best Local Similarity 100.0%; Pred. No. 0.16;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 PVTAESEPKKP 10
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 DB 197 pvtasepkkp 206

Search completed: October 12, 2002, 20:43:41
 Job time: 13936 sec

GenCore version 4.5
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OM protein - protein search, using sw model

Run on: October 12, 2002, 16:54:45 ; Search time 32.92 Seconds
(without alignments)
7.420 Million cell updates/sec

Title: US-09-408-578A-1
Perfect score: 52
Sequence: 1 PVVAESPCKP 10

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 231628 seqs, 24425594 residues

Total number of hits satisfying chosen parameters: 231628

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : Issued Patents AA:*
1: /cgn2_6/ptodata/2/1aa/5A.COMB.pep:*
2: /cgn2_6/ptodata/2/1aa/5B.COMB.pep:*
3: /cgn2_6/ptodata/2/1aa/6A.COMB.pep:*
4: /cgn2_6/ptodata/2/1aa/6B.COMB.pep:*
5: /cgn2_6/ptodata/2/1aa/6C.COMB.pep:*
6: /cgn2_6/ptodata/2/1aa/backfile1.pep:*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	52	100.0	194	4 US-09-364-083-2	Sequence 2, App1
2	52	100.0	207	4 US-08-235-836C-36	Sequence 36, App1
3	52	100.0	209	4 US-08-235-836C-32	Sequence 32, App1
4	52	100.0	210	1 US-08-158-353-3	Sequence 3, App1
5	52	100.0	210	4 US-08-209-603E-15	Sequence 15, App1
6	52	100.0	210	4 US-08-235-836C-30	Sequence 30, App1
7	52	100.0	212	1 US-08-158-353-4	Sequence 4, App1
8	52	100.0	212	4 US-09-196-293-11	Sequence 11, App1
9	52	100.0	212	4 US-08-209-603E-11	Sequence 11, App1
10	52	100.0	212	4 US-08-235-836C-34	Sequence 34, App1
11	52	100.0	466	4 US-08-235-836C-107	Sequence 107, App
12	52	100.0	588	4 US-08-235-836C-110	Sequence 110, App
13	52	100.0	588	4 US-08-235-836C-122	Sequence 122, App
14	49	94.2	212	1 US-08-031-295-2	Sequence 2, App1
15	49	94.2	212	1 US-08-158-353-2	Sequence 2, App1
16	49	94.2	212	4 US-07-903-80-2	Sequence 2, App1
17	40	76.9	209	4 US-09-196-293-15	Sequence 15, App1
18	35	67.3	1255	3 US-08-947-823-3	Sequence 3, App1
19	35	67.3	1257	3 US-08-947-823-5	Sequence 5, App1
20	34	65.4	984	1 US-08-242-932-2	Sequence 2, App1
21	34	65.4	984	1 US-08-714-481-2	Sequence 2, App1
22	34	65.4	984	5 PCT-US95-06111-2	Sequence 2, App1
23	34	65.4	1098	4 US-08-923-992A-8	Sequence 8, App1
24	34	65.4	1104	4 US-08-923-992A-4	Sequence 4, App1
25	34	65.4	1128	4 US-08-923-992A-6	Sequence 6, App1
26	34	65.4	1164	4 US-08-923-992A-2	Sequence 2, App1
27	34	65.4	1164	4 US-08-923-992A-10	Sequence 10, App1

28	33	63.5	22	4 US-08-557-006C-34	Sequence 34, App1
29	33	63.5	197	3 US-08-415-655-6	Sequence 6, App1
30	33	63.5	417	3 US-08-705-771-18	Sequence 18, App1
31	33	63.5	492	4 US-09-342-749-2	Sequence 2, App1
32	33	63.5	552	4 US-08-557-006C-40	Sequence 40, App1
33	33	63.5	593	3 US-09-000-145-4	Sequence 4, App1
34	33	63.5	933	3 US-08-293-728-2	Sequence 2, App1
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36	32	61.5	164	1 US-08-357-125-4	Sequence 4, App1
37	32	61.5	165	5 PCT-US95-03666-4	Sequence 4, App1
38	32	61.5	273	1 US-08-220-379B-6	Sequence 6, App1
39	32	61.5	273	1 US-08-341-456A-11	Sequence 11, App1
40	32	61.5	273	3 US-08-478-414A-11	Sequence 11, App1
41	32	61.5	273	3 US-08-325-240A-11	Sequence 11, App1
42	32	61.5	273	4 US-08-898-982-11	Sequence 11, App1
43	32	61.5	273	4 US-08-482-918-55	Sequence 55, App1
44	32	61.5	273	4 US-09-224-661-55	Sequence 55, App1
45	32	61.5	273	4 US-08-336-728A-55	Sequence 55, App1

ALIGNMENTS

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RESULT 1
; Sequence 2, Application US/09364083
; Patent No. 6210676
; GENERAL INFORMATION:
; APPLICANT: Callister, Steven M
; APPLICANT: Lovitch, Steven D
; APPLICANT: Schell, Ronald F
; APPLICANT: Jobe, Dean A
; TITLE OF INVENTION: Compositions and Method using the Borrellia
; TITLE OF INVENTION: Protein C (ospc) for the diagnosis and prevention of
; TITLE OF INVENTION: Lyme Disease
; FILE REFERENCE: B. burgdorferi OspC
; CURRENT APPLICATION NUMBER: US/09/364,083
; EARLIER FILING DATE: 1999-07-30
; EARLIER APPLICATION NUMBER: 60/094,955
; SOFTWARE: Patentln Ver. 2.0
; SEQ ID NO 2
; LENGTH: 194
; TYPE: PRT
; ORGANISM: Borrellia burgdorferi
US-09-364-083-2
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Query Match      100.0%  Score 52;  DB 4;  Length 194;
Best Local Similarity 100.0%;  Pred. NO. 0.029;
Matches 10;  Conservative 0;  Mismatches 0;  Indels 0;  Gaps 0;
QY      1 PVVAESPCKP 10
Db      185 PVVAESPCKP 194
-----
RESULT 2
US-08-235-836C-36
; Sequence 36, Application US/08235836C
; Patent No. 6248562
; GENERAL INFORMATION:
; APPLICANT: Dunn, John J.
; APPLICANT: Luft, Benjamin J.
; TITLE OF INVENTION: No. 6248562el Chimeric Proteins Comprising
; TITLE OF INVENTION: Borrellia Polypeptides and Uses Therefor
; NUMBER OF SEQUENCES: 144
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Brookhaven National Laboratory
; STREET:
; CITY: Upton
```

STATE: NY
COUNTRY: USA
ZIP: 11973
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/235,836C
FILING DATE: 29-APR-1994
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/148,191
FILING DATE: 01-11-93
ATTORNEY/AGENT INFORMATION:
NAME: Bogosian, Margaret C.
REGISTRATION NUMBER: 25,324
REFERENCE/DOCKET NUMBER: BNL93-28A
TELECOMMUNICATION INFORMATION:
TELEPHONE: (516) 282-7338
TELEFAX: (516) 282-3729
INFORMATION FOR SEQ ID NO: 36:
SEQUENCE CHARACTERISTICS:
LENGTH: 207 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-235-836C-36

Query Match 100.0%; Score 52; DB 4; Length 207;
Best Local Similarity 100.0%; Pred. No. 0.031;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 PVVAESPCKP 10
DB 198 PVVAESPCKP 207

RESULT 3
US-08-235-836C-32
Sequence 32, Application US/08235836C
Patent No. 6248562
GENERAL INFORMATION:
APPLICANT: Dunn, John J.
APPLICANT: Luft, Benjamin J.
TITLE OF INVENTION: No. 6248562el Chimeric Proteins Comprising
TITLE OF INVENTION: Borrella Polypeptides and Uses Therefor
NUMBER OF SEQUENCES: 144
CORRESPONDENCE ADDRESS:
ADDRESSEE: Brookhaven National Laboratory
STREET:
CITY: Upton
STATE: NY
COUNTRY: USA
ZIP: 11973
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/235,836C
FILING DATE: 29-APR-1994
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/148,191
FILING DATE: 01-11-93
ATTORNEY/AGENT INFORMATION:
NAME: Bogosian, Margaret C.
REGISTRATION NUMBER: 25,324
REFERENCE/DOCKET NUMBER: BNL93-28A

TELECOMMUNICATION INFORMATION:
TELEPHONE: (516) 282-7338
TELEFAX: (516) 282-3729
INFORMATION FOR SEQ ID NO: 32:
SEQUENCE CHARACTERISTICS:
LENGTH: 209 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-235-836C-32

Query Match 100.0%; Score 52; DB 4; Length 209;
Best Local Similarity 100.0%; Pred. No. 0.032;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 PVVAESPCKP 10
DB 200 PVVAESPCKP 209

RESULT 4
US-08-158-353-3
Sequence 3, Application US/08158353
Patent No. 5620862
GENERAL INFORMATION:
APPLICANT: Padula, Steven J.
TITLE OF INVENTION: Methods for Diagnosing Early Lyme
TITLE OF INVENTION: Disease
NUMBER OF SEQUENCES: 7
CORRESPONDENCE ADDRESS:
ADDRESSEE: Hamilton, Brook, Smith & Reynolds, P.C.
STREET: Two Millitia Drive
CITY: Lexington
STATE: MA
COUNTRY: USA
ZIP: 02173
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/158,353
FILING DATE:
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Carroll, Alice O
REGISTRATION NUMBER: 33,542
REFERENCE/DOCKET NUMBER: DCT93-05
TELECOMMUNICATION INFORMATION:
TELEPHONE: 617-861-6240
TELEFAX: 617-861-9540
INFORMATION FOR SEQ ID NO: 3:
SEQUENCE CHARACTERISTICS:
LENGTH: 210 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-158-353-3

Query Match 100.0%; Score 52; DB 1; Length 210;
Best Local Similarity 100.0%; Pred. No. 0.032;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 PVVAESPCKP 10
DB 201 PVVAESPCKP 210

RESULT 5

US-08-209-603E-15
; Sequence 15, Application US/08209603E
; Patent No. 6248538
; GENERAL INFORMATION:
; APPLICANT: FUCHS, RENATE
; APPLICANT: MILSK, BETTINA
; APPLICANT: PREAC-MORSIC, VERA
; APPLICANT: MOTZ, MANFRED
; APPLICANT: SOUTSCHECK, ERMIN
; TITLE OF INVENTION: IMMUNOLOGICALLY ACTIVE PROTEINS
; TITLE OF INVENTION: FROM BORRELLA BURGDOFFERI
; NUMBER OF SEQUENCES: 15
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: BROOKS HAIDT HAFNER & DELAHUNTY
; STREET: 99 PARK AVENUE
; CITY: NEW YORK
; STATE: NY
; COUNTRY: USA
; ZIP: 10016
; COMPUTER READABLE FORM:
; MEDIUM TYPE: 3.5" FLOPPY DISC
; COMPUTER: AT&T - IBM COMPATIBLE
; OPERATING SYSTEM: MS-DOS Version 6.2
; SOFTWARE: ASCII
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/209,603E
; FILING DATE: 10-MAR-1994
; CLASSIFICATION: 436
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: PCT/EP90/02282
; FILING DATE: 21-DEC-1990
; APPLICATION NUMBER: US 07/862,535
; FILING DATE: 19-JUN-1992
; ATTORNEY/AGENT INFORMATION:
; NAME: ROBINSON, WILLIAM R.
; REGISTRATION NUMBER: 27,224
; REFERENCE/DOCKET NUMBER: LKR-9217-A
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (212) 697-3355
; TELEFAX: (212) 557-5635
; INFORMATION FOR SEQ ID NO: 15:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 210
; TYPE: AMINO ACID
; TOPOLOGY: LINEAR
; MOLECULE TYPE: PROTEIN
; DESCRIPTION: N/A
; HYPOTHEICAL: N/A
; ANTI-SENSE: N/A
; FRAGMENT TYPE: N/A
; ORGANISM: B. BURGDOFFERI
; IMMEDIATE SOURCE:
; LIBRARY: DSM 5662
; POSITION IN GENOME: N/A
; FEATURE:
; IDENTIFICATION METHOD: amino acid analysis
; PUBLICATION INFORMATION: N/A
; US-08-209-603E-15

Query Match 100.0%; Score 52; DB 4; Length 210;
Best Local Similarity 100.0%; Pred. No. 0.032;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 PVVAESPKP 10
| | | | | | | | | |
Db 201 PVVAESPKP 210

RESULT 6
US-08-235-836C-30
; Sequence 30, Application US/08235836C

Patent No. 6248562
; GENERAL INFORMATION:
; APPLICANT: Dunn, John J.
; APPLICANT: Luft, Benjamin J.
; TITLE OF INVENTION: No. 6248562a1 Chimeric Proteins Comprising
; TITLE OF INVENTION: Borrella Polypeptides and uses therefor
; NUMBER OF SEQUENCES: 144
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Brookhaven National Laboratory
; STREET:
; CITY: Upton
; STATE: NY
; COUNTRY: USA
; ZIP: 11973
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/235,836C
; FILING DATE: 29-APR-1994
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/148,191
; FILING DATE: 01-11-93
; ATTORNEY/AGENT INFORMATION:
; NAME: Bogostian, Margaret C.
; REGISTRATION NUMBER: 25,324
; REFERENCE/DOCKET NUMBER: BNL93-28A
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (516) 282-7338
; TELEFAX: (516) 282-3729
; INFORMATION FOR SEQ ID NO: 30:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 210 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; US-08-235-836C-30

Query Match 100.0%; Score 52; DB 4; Length 210;
Best Local Similarity 100.0%; Pred. No. 0.032;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 PVVAESPKP 10
| | | | | | | | | |
Db 201 PVVAESPKP 210

RESULT 7
US-08-158-353-4
; Sequence 4, Application US/08158353
; Patent No. 5620862
; GENERAL INFORMATION:
; APPLICANT: Padula, Steven J.
; TITLE OF INVENTION: Methods for Diagnosing Early Lyme
; TITLE OF INVENTION: Disease
; NUMBER OF SEQUENCES: 7
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Hamilton, Brook, Smith & Reynolds, P.C.
; STREET: Two Militia Drive
; CITY: Lexington
; STATE: MA
; COUNTRY: USA
; ZIP: 02173
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/158,353
FILING DATE:
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Carroll, Alice O.
REGISTRATION NUMBER: 33,542
REFERENCE/DOCKET NUMBER: UCT93-05
TELECOMMUNICATION INFORMATION:
TELEPHONE: 617-861-6240
TELEFAX: 617-861-9540
INFORMATION FOR SEQ ID NO: 4:
SEQUENCE CHARACTERISTICS:
LENGTH: 212 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-158-353-4

Query Match 100.0%; Score 52; DB 1; Length 212;
Best Local Similarity 100.0%; Pred. No. 0.032;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 PVVAESPKP 10
|||||
DB 203 PVVAESPKP 212

RESULT 8
US-09-196-293-11
Sequence 11, Application US/09196293
Patent No. 6183755
GENERAL INFORMATION:
APPLICANT: Fuchs, Renate
APPLICANT: Motz, Manfred
APPLICANT: Soutscheck, Erwin
APPLICANT: Wilske, Bettina
APPLICANT: Preac-Mursic, Vera
TITLE OF INVENTION: Active proteins from Borrelia
FILE REFERENCE: 738,001US2
CURRENT APPLICATION NUMBER: US/09/196,293
CURRENT FILING DATE: 1998-11-19
EARLIER APPLICATION NUMBER: US 08/209,603
EARLIER FILING DATE: 1994-03-10
EARLIER APPLICATION NUMBER: US 07/862,535
EARLIER FILING DATE: 1992-06-19
EARLIER APPLICATION NUMBER: WO PCT/EP90/02282
EARLIER FILING DATE: 1990-12-21
EARLIER APPLICATION NUMBER: DE P39 42,728.5
EARLIER FILING DATE: 1989-12-22
EARLIER APPLICATION NUMBER: DE P40 18 988.0
EARLIER FILING DATE: 1990-06-13
NUMBER OF SEQ ID NOS: 16
SOFTWARE: FastSeq for Windows Version 4.0
SEQ ID NO 11
LENGTH: 212
TYPE: PRT
ORGANISM: Borrelia burgdorferi
US-09-196-293-11

Query Match 100.0%; Score 52; DB 4; Length 212;
Best Local Similarity 100.0%; Pred. No. 0.032;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 PVVAESPKP 10
|||||
DB 203 PVVAESPKP 212

RESULT 9

US-08-209-603E-11
Sequence 11, Application US/08209603E
Patent No. 6248538
GENERAL INFORMATION:
APPLICANT: FUCHS, RENATE
APPLICANT: WILSKE, BETTINA
APPLICANT: PREAC-MURISIC, VERA
APPLICANT: MOTZ, MANFRED
APPLICANT: SOUTSCHECK, ERWIN
TITLE OF INVENTION: IMMUNOLOGICALLY ACTIVE PROTEINS
NUMBER OF SEQUENCES: 15
CORRESPONDENCE ADDRESS:
ADDRESSEE: BROOKS HAIDT HAFNER & DELAHUNTY
STREET: 99 PARK AVENUE
CITY: NEW YORK
STATE: NY
COUNTRY: USA
ZIP: 10016

COMPUTER READABLE FORM:
MEDIUM TYPE: 3.5" FLOPPY DISC
COMPUTER: AT&T - IBM COMPATIBLE
OPERATING SYSTEM: MS-DOS Version 6.2
SOFTWARE: ASCII
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/209,603E
FILING DATE: 10-MAR-1994
CLASSIFICATION: 436
PRIOR APPLICATION DATA:
APPLICATION NUMBER: PCT/EP90/02282
FILING DATE: 21-DEC-1990
APPLICATION NUMBER: US 07/862,535
FILING DATE: 19-JUN-1992
ATTORNEY/AGENT INFORMATION:
NAME: ROBINSON, WILLIAM R.
REGISTRATION NUMBER: 27,224
REFERENCE/DOCKET NUMBER: LKR-9217-A
TELECOMMUNICATION INFORMATION:
TELEPHONE: (212) 557-5635
TELEFAX: (212) 557-5635
INFORMATION FOR SEQ ID NO: 11:
SEQUENCE CHARACTERISTICS:
LENGTH: 212
TYPE: AMINO ACID
TOPOLOGY: LINEAR
MOLECULE TYPE: PROTEIN
DESCRIPTION: N/A
HYPOTHETICAL: N/A
ANTI-SENSE: N/A
FRAGMENT TYPE: N/A
ORIGINAL SOURCE:
ORGANISM: B. BURGDOFFERI
IMMEDIATE SOURCE:
LIBRARY: DSM 5662
POSITION IN GENOME: N/A
FEATURE:
IDENTIFICATION METHOD: amino acid analysis
PUBLICATION INFORMATION: N/A
ANTI-SENSE: N/A
FRAGMENT TYPE: INTERNAL
ORIGINAL SOURCE:

US-08-209-603E-11

Query Match 100.0%; Score 52; DB 4; Length 212;
Best Local Similarity 100.0%; Pred. No. 0.032;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 PVVAESPKP 10
|||||
DB 203 PVVAESPKP 212

```
RESULT 10
US-08-235-836C-34
; Sequence 34, Application US/08235836C
; Patent No. 6248562
; GENERAL INFORMATION:
; APPLICANT: Dunn, John J.
; TITLE OF INVENTION: No. 6248562el Chimeric Proteins Comprising
; NUMBER OF SEQUENCES: 144
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Brookhaven National Laboratory
; STREET:
; CITY: Upton
; STATE: NY
; COUNTRY: USA
; ZIP: 11973
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/235,836C
; FILING DATE: 29-Apr-1994
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/148,191
; FILING DATE: 01-11-93
; ATTORNEY/AGENT INFORMATION:
; NAME: Bogosian, Margaret C.
; REGISTRATION NUMBER: 25,324
; REFERENCE/DOCKET NUMBER: BNL93-28A
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (516) 282-7338
; TELEFAX: (516) 282-3729
; INFORMATION FOR SEQ ID NO: 34:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 212 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; US-08-235-836C-34

Query Match          100.0%; Score 52; DB 4; Length 212;
Best Local Similarity 100.0%; Pred. No. 0.032;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 PYVAESPKRP 10
       |||||
Db      203 PYVAESPKRP 212

RESULT 11
US-08-235-836C-107
; Sequence 107, Application US/08235836C
; Patent No. 6248562
; GENERAL INFORMATION:
; APPLICANT: Dunn, John J.
; TITLE OF INVENTION: No. 6248562el Chimeric Proteins Comprising
; NUMBER OF SEQUENCES: 144
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Brookhaven National Laboratory
; STREET:
; CITY: Upton
; STATE: NY
; COUNTRY: USA
; ZIP: 11973
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
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COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/235,836C
; FILING DATE: 29-Apr-1994
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/148,191
; FILING DATE: 01-11-93
; ATTORNEY/AGENT INFORMATION:
; NAME: Bogosian, Margaret C.
; REGISTRATION NUMBER: 25,324
; REFERENCE/DOCKET NUMBER: BNL93-28A
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (516) 282-7338
; TELEFAX: (516) 282-3729
; INFORMATION FOR SEQ ID NO: 107:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 466 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; US-08-235-836C-107

Query Match          100.0%; Score 52; DB 4; Length 466;
Best Local Similarity 100.0%; Pred. No. 0.075;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY      1 PYVAESPKRP 10
       |||||
Db      457 PYVAESPKRP 466

RESULT 12
US-08-235-836C-110
; Sequence 110, Application US/08235836C
; Patent No. 6248562
; GENERAL INFORMATION:
; APPLICANT: Dunn, John J.
; TITLE OF INVENTION: No. 6248562el Chimeric Proteins Comprising
; NUMBER OF SEQUENCES: 144
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Brookhaven National Laboratory
; STREET:
; CITY: Upton
; STATE: NY
; COUNTRY: USA
; ZIP: 11973
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/235,836C
; FILING DATE: 29-Apr-1994
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/148,191
; FILING DATE: 01-11-93
; ATTORNEY/AGENT INFORMATION:
; NAME: Bogosian, Margaret C.
; REGISTRATION NUMBER: 25,324
; REFERENCE/DOCKET NUMBER: BNL93-28A
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (516) 282-7338
; TELEFAX: (516) 282-3729
; INFORMATION FOR SEQ ID NO: 110:
; SEQUENCE CHARACTERISTICS:
```

LENGTH: 466 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-235-836C-110

Query Match
Best Local Similarity 100.0%; Score 52; DB 4; Length 466;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 PVVAESPKKP 10
|||||
DB 201 PVVAESPKKP 210

RESULT 13
US-08-235-836C-122
Sequence 122, Application US/08235836C
Patent No. 6248562
GENERAL INFORMATION:
APPLICANT: Dunn, John J.
APPLICANT: Luft, Benjamin J.
TITLE OF INVENTION: No. 6248562el Chimeric Proteins Comprising
TITLE OF INVENTION: Borrelia Polypeptides and Uses Therefor
NUMBER OF SEQUENCES: 144
CORRESPONDENCE ADDRESS:
ADDRESSEE: Brookhaven National Laboratory
STREET:
CITY: Upton
STATE: NY
COUNTRY: USA
ZIP: 11973
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/235.836C
FILING DATE: 29-APR-1994
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/148,191
FILING DATE: 01-11-93
ATTORNEY/AGENT INFORMATION:
NAME: Bogosian, Margaret C.
REGISTRATION NUMBER: 25,324
REFERENCE/DOCKET NUMBER: BNL93-28A
TELEPHONE: (516) 282-7338
TELEFAX: (516) 282-3729
INFORMATION FOR SEQ ID NO: 122:
SEQUENCE CHARACTERISTICS:
LENGTH: 588 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-235-836C-122

Query Match
Best Local Similarity 100.0%; Score 52; DB 4; Length 588;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 PVVAESPKKP 10
|||||
DB 467 PVVAESPKKP 476

RESULT 14
US-08-031-295-2
Sequence 2, Application US/08031295

Patent No. 5530103
GENERAL INFORMATION:
APPLICANT: LIVER, Ian
APPLICANT: DORNER, Friedrich
TITLE OF INVENTION: METHOD AND COMPOSITION FOR THE
TITLE OF INVENTION: PREVENTION OF LYME DISEASE
NUMBER OF SEQUENCES: 3
CORRESPONDENCE ADDRESS:
ADDRESSEE: Foley & Lardner
STREET: 3000 K Street, N.W., Suite 500
CITY: Washington, D.C.
COUNTRY: USA
ZIP: 20007-5109
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/031.295
FILING DATE: 19930312
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/903,580
FILING DATE: 25-JUN-1992
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/824,161
FILING DATE: 22-JAN-1992
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/727,245
FILING DATE: 11-JUL-1991
ATTORNEY/AGENT INFORMATION:
NAME: BENT, Stephen A.
REGISTRATION NUMBER: 29,768
REFERENCE/DOCKET NUMBER: 30472/142 IMMU
TELEPHONE: (202)672-5300
TELEFAX: (202)672-5399
TELEX: 904136
INFORMATION FOR SEQ ID NO: 2:
SEQUENCE CHARACTERISTICS:
LENGTH: 212 amino acids
TYPE: AMINO ACID
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-031-295-2

Query Match
Best Local Similarity 94.2%; Score 49; DB 1; Length 212;
Matches 9; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 PVVAETPKKP 10
|||||
DB 203 PVVAETPKKP 212

RESULT 15
US-08-158-353-2
Sequence 2, Application US/08158353
Patent No. 5620862
GENERAL INFORMATION:
APPLICANT: Padula, Steven J.
TITLE OF INVENTION: Methods for Diagnosing Early Lyme
TITLE OF INVENTION: Disease
NUMBER OF SEQUENCES: 7
CORRESPONDENCE ADDRESS:
ADDRESSEE: Hamilton, Brook, Smith & Reynolds, P.C.
STREET: Two Millitia Drive
CITY: Lexington
STATE: MA
COUNTRY: USA
ZIP: 02173

COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/158,353
FILING DATE:
CLASSIFICATION: 435
ATTORNEY/AGENT INFORMATION:
NAME: Carroll, Alice O.
REGISTRATION NUMBER: 33,542
REFERENCE/DOCKET NUMBER: UCT93-05
TELECOMMUNICATION INFORMATION:
TELEPHONE: 617-861-6240
TELEFAX: 617-861-9540
INFORMATION FOR SEQ ID NO: 2:
SEQUENCE CHARACTERISTICS:
LENGTH: 212 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-158-353-2

Query Match 94.2%; Score 49; DB 1; Length 212;
Best Local Similarity 90.0%; Pred. No. 0.11;
Matches 9; Conservative 1; Mismatches 0; Indels 0; Gaps 0;
QY 1 PYVAESPCKP 10
|||||:||||
Db 203 PYVAENPKKP 212

Search completed: October 12, 2002, 20:44:28
Job time: 13783 sec

GenCore version 4.5
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OM protein - protein search, using sw model

Run on: October 12, 2002, 17:08:10 ; Search time 49.13 Seconds

(without alignments)
19,558 Million cell updates/sec

Title: US-09-408-578A-1

Sequence: 1 PVVAESPCKP 10

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 283138 seqs, 96089334 residues

Total number of hits satisfying chosen parameters: 283138

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%

Listing first 45 summaries

Database :

1: PIR.71.*
2: PIR1.*
3: PIR3.*
4: PIR4.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	52	100.0	190	2 S70273	outer surface prot
2	52	100.0	191	2 S70278	outer surface prot
3	52	100.0	191	2 S70284	outer surface prot
4	52	100.0	193	2 S70265	outer surface prot
5	52	100.0	193	2 S70274	outer surface prot
6	52	100.0	193	2 S70279	outer surface prot
7	52	100.0	194	2 S70268	outer surface prot
8	52	100.0	194	2 S70277	outer surface prot
9	52	100.0	207	2 I40271	outer surface prot
10	52	100.0	207	2 I40276	outer surface prot
11	52	100.0	207	2 S69919	outer surface prot
12	52	100.0	207	2 S69924	outer surface prot
13	52	100.0	207	2 S37727	outer surface prot
14	52	100.0	209	2 I40273	outer surface prot
15	52	100.0	209	2 I40270	outer surface prot
16	52	100.0	209	2 S69926	outer surface prot
17	52	100.0	209	2 I40281	outer surface prot
18	52	100.0	209	2 I40285	outer surface prot
19	52	100.0	209	2 S69917	outer surface prot
20	52	100.0	210	2 I40272	outer surface prot
21	52	100.0	210	2 S69920	outer surface prot
22	52	100.0	210	2 S69925	outer surface prot
23	52	100.0	210	2 I40280	outer surface prot
24	52	100.0	210	2 I40284	outer surface prot
25	52	100.0	210	2 C70218	outer surface prot
26	52	100.0	210	2 I40144	outer surface prot
27	52	100.0	210	2 S69927	outer surface prot
28	52	100.0	210	2 S69923	outer surface prot
29	52	100.0	211	2 I40277	outer surface prot

30	52	100.0	211	2 I40278	outer surface prot
31	52	100.0	211	2 I40282	outer surface prot
32	52	100.0	211	2 I40145	outer surface prot
33	52	100.0	211	2 I40268	outer surface prot
34	52	100.0	211	2 S69930	outer surface prot
35	52	100.0	211	2 S69918	outer surface prot
36	52	100.0	211	2 S69932	outer surface prot
37	52	100.0	211	2 S69928	outer surface prot
38	52	100.0	211	2 S69929	outer surface prot
39	52	100.0	212	2 I40279	outer surface prot
40	52	100.0	212	2 S69921	outer surface prot
41	52	100.0	212	2 S69922	outer surface prot
42	52	100.0	212	2 I40275	outer surface prot
43	52	100.0	212	2 S20543	outer surface prot
44	52	100.0	214	2 S69916	outer surface prot
45	51	98.1	191	2 I40153	outer surface prot

ALIGNMENTS

RESULT 1
S70273
outer surface protein C - Lyme disease spirochete
C:Species: Borrelia burgdorferi (Lyme disease spirochete)
C:Date: 12-Feb-1998 #sequence_revision 20-Feb-1998 #text_change 26-May-2000
C:Accession: S70273
R:Livey, I.; Gibbs, C.P.; Schuster, R.; Dörner, F.
Mol. Microbiol. 18, 257-269, 1995
A>Title: Evidence for lateral transfer and recombination in OspC variation in Lyme d
A:Reference number: S70255; MUID:96296448
A:Accession: S70273
A>Status: nucleic acid sequence not shown
A:Molecule type: DNA
A:Residues: 1-190 <Lit>
A:Cross-references: EMBL:L42870; NID:9858737; PIDN:AAB37013.1; PID:91695228
A:Experimental source: strain VSDA
C:Genetics:
A:Gene: ospC
C:Superfamily: Lyme disease spirochete surface protein C

Query Match 100.0%; Score 52; DB 2; Length 190;
Best Local Similarity 100.0%; Pred. No. 0.036;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 PVVAESPCKP 10
DB 181 PVVAESPCKP 190

RESULT 2
S70278
outer surface protein C - Lyme disease spirochete
C:Species: Borrelia burgdorferi (Lyme disease spirochete)
C:Date: 12-Feb-1998 #sequence_revision 20-Feb-1998 #text_change 26-May-2000
C:Accession: S70278
R:Livey, I.; Gibbs, C.P.; Schuster, R.; Dörner, F.
Mol. Microbiol. 18, 257-269, 1995
A>Title: Evidence for lateral transfer and recombination in OspC variation in Lyme d
A:Reference number: S70255; MUID:96296448
A:Accession: S70278
A>Status: nucleic acid sequence not shown
A:Molecule type: DNA
A:Residues: 1-191 <Lit>
A:Cross-references: EMBL:L42871; NID:9858738; PIDN:AAB37014.1; PID:91695229
A:Experimental source: strain VS461
C:Genetics:
A:Gene: ospC
C:Superfamily: Lyme disease spirochete surface protein C

Query Match 100.0%; Score 52; DB 2; Length 191;

Best Local Similarity 100.0%; Score 52; DB 2; Length 193;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 PVVAESPCKP 10
|||||
Db 182 PVVAESPCKP 191

RESULT 3

S70284

outer surface protein C - Lyme disease spirochete

C:Species: Borrelia burgdorferi (Lyme disease spirochete)

C>Date: 12-Feb-1998 #sequence_rev1sion 20-Feb-1998 #text_change 26-May-2000

C:Accession: S70284

R:Livey, I.; Gibbs, C.P.; Schuster, R.; Dorner, F.

Mol. Microbiol. 18, 257-269, 1995

A:Title: Evidence for lateral transfer and recombination in OspC variation in Lyme disease

A:Reference number: S70255; PMID:96296448

A:Accession: S70284

A>Status: nucleic acid sequence not shown

A:Molecule type: DNA

A:Residues: 1-191 <Lit>

A:Cross-references: EMBL:L42896; NID:g858724; PIDN:AAB37004.1; PID:g1695221

A:Experimental source: strain 27579

C:Genetics:

A:Gene: ospC

C:Superfamily: Lyme disease spirochete surface protein C

Query Match

Best Local Similarity 100.0%; Score 52; DB 2; Length 191;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 PVVAESPCKP 10
|||||
Db 182 PVVAESPCKP 191

RESULT 4

S70265

outer surface protein C - Lyme disease spirochete

C:Species: Borrelia burgdorferi (Lyme disease spirochete)

C>Date: 12-Feb-1998 #sequence_rev1sion 20-Feb-1998 #text_change 26-May-2000

C:Accession: S70265

R:Livey, I.; Gibbs, C.P.; Schuster, R.; Dorner, F.

Mol. Microbiol. 18, 257-269, 1995

A:Title: Evidence for lateral transfer and recombination in OspC variation in Lyme disease

A:Reference number: S70255; PMID:96296448

A:Accession: S70265

A>Status: nucleic acid sequence not shown

A:Molecule type: DNA

A:Residues: 1-193 <Lit>

A:Cross-references: EMBL:L42884; NID:g858710; PIDN:AAB36992.1; PID:g1695210

A:Experimental source: strain J1

C:Genetics:

A:Gene: ospC

C:Superfamily: Lyme disease spirochete surface protein C

Query Match
Best Local Similarity 100.0%; Score 52; DB 2; Length 193;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 PVVAESPCKP 10
|||||
Db 184 PVVAESPCKP 193

RESULT 5

S70274

outer surface protein C - Lyme disease spirochete

C:Species: Borrelia burgdorferi (Lyme disease spirochete)

C>Date: 12-Feb-1998 #sequence_rev1sion 20-Feb-1998 #text_change 26-May-2000

C:Accession: S70274

R:Livey, I.; Gibbs, C.P.; Schuster, R.; Dorner, F.

Mol. Microbiol. 18, 257-269, 1995

A:Title: Evidence for lateral transfer and recombination in OspC variation in Lyme

A:Reference number: S70255; PMID:96296448

A:Accession: S70274

A>Status: nucleic acid sequence not shown

A:Molecule type: DNA

A:Residues: 1-193 <Lit>

A:Cross-references: EMBL:L42892; NID:g858720; PIDN:AAB37000.1; PID:g1695217

A:Experimental source: strain acal

C:Genetics:

A:Gene: ospC

C:Superfamily: Lyme disease spirochete surface protein C

Query Match
Best Local Similarity 100.0%; Score 52; DB 2; Length 193;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 PVVAESPCKP 10
|||||
Db 184 PVVAESPCKP 193

RESULT 6

S70279

outer surface protein C - Lyme disease spirochete

C:Species: Borrelia burgdorferi (Lyme disease spirochete)

C>Date: 12-Feb-1998 #sequence_rev1sion 20-Feb-1998 #text_change 26-May-2000

C:Accession: S70279

R:Livey, I.; Gibbs, C.P.; Schuster, R.; Dorner, F.

Mol. Microbiol. 18, 257-269, 1995

A:Title: Evidence for lateral transfer and recombination in OspC variation in Lyme

A:Reference number: S70255; PMID:96296448

A:Accession: S70279

A>Status: nucleic acid sequence not shown

A:Molecule type: DNA

A:Residues: 1-193 <Lit>

A:Cross-references: EMBL:L42898; NID:g858729; PIDN:AAB37007.1; PID:g1695223

A:Experimental source: strain 25015

C:Genetics:

A:Gene: ospC

C:Superfamily: Lyme disease spirochete surface protein C

Query Match
Best Local Similarity 100.0%; Score 52; DB 2; Length 193;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 PVVAESPCKP 10
|||||
Db 184 PVVAESPCKP 193

RESULT 7

S70268

outer surface protein C - Lyme disease spirochete

C:Species: Borrelia burgdorferi (Lyme disease spirochete)

C>Date: 12-Feb-1998 #sequence_rev1sion 20-Feb-1998 #text_change 26-May-2000

C:Accession: S70268

R:Livey, I.; Gibbs, C.P.; Schuster, R.; Dorner, F.

Mol. Microbiol. 18, 257-269, 1995

A:Title: Evidence for lateral transfer and recombination in OspC variation in Lyme

A:Reference number: S70255; PMID:96296448

A:Accession: S70268

A>Status: nucleic acid sequence not shown

A:Molecule type: DNA

A:Residues: 1-194 <Lit>

A:Cross-references: EMBL:L42888; NID:g858716; PIDN:AAB36996.1; PID:g1695213

A:Experimental source: strain H9

C:Genetics:

A:Gene: ospC

C:Superfamily: Lyme disease spirochete surface protein C

Query Match
Best Local Similarity 100.0%; Score 52; DB 2; Length 194;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 PVVAESPCKP 10
|||||
Db 185 PVVAESPCKP 194

RESULT 8
S70277

outer surface protein C - Lyme disease spirochete
C:Species: Borrelia burgdorferi (Lyme disease spirochete)
C:Date: 12-Feb-1998 #sequence_revision 20-Feb-1998 #text_change 26-May-2000
C:Accession: S70277
R:Ulevy, I.; Gibbs, C.P.; Schuster, R.; Dörner, F.
Mol. Microbiol. 18, 257-269, 1995
A:Title: Evidence for lateral transfer and recombination in OspC variation in Lyme disease
A:Reference number: S70255; MUID:96296448
A:Accession: S70277
A:Status: nucleic acid sequence not shown
A:Molecule type: DNA
A:Residues: 1-194 <LIV>
A:Cross-references: EMBL:LA2873; NID:9858740; PIDN:AA837016.1; PID:91695231
A:Experimental source: strain SIMON
C:Genetics:
A:Gene: ospC
C:Superfamily: Lyme disease spirochete surface protein C

Query Match
Best Local Similarity 100.0%; Score 52; DB 2; Length 194;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 PVVAESPCKP 10
|||||
Db 185 PVVAESPCKP 194

RESULT 9
I40271

outer surface protein C precursor - Borrelia garinii
C:Species: Borrelia garinii
C:Date: 04-Sep-1997 #sequence_revision 04-Sep-1997 #text_change 26-May-2000
C:Accession: I40271
R:Fukunaga, M.; Hamase, A.
J. Clin. Microbiol. 33, 2415-2420, 1995
A:Title: Outer surface protein C gene sequence analysis of Borrelia burgdorferi sensu lato
A:Reference number: I40269; MUID:96025162
A:Accession: I40271
A:Status: preliminary; translated from GB/EMBL/DBDJ
A:Molecule type: DNA
A:Residues: 1-207 <RES>
A:Cross-references: GB:D49377; NID:G1041103; PIDN:BA08375.1; PID:G1041104
C:Superfamily: Lyme disease spirochete surface protein C

Query Match
Best Local Similarity 100.0%; Score 52; DB 2; Length 207;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 PVVAESPCKP 10
|||||
Db 198 PVVAESPCKP 207

RESULT 10
I40276

outer surface protein C precursor - Borrelia garinii
C:Species: Borrelia garinii

C:Date: 04-Sep-1997 #sequence_revision 04-Sep-1997 #text_change 26-May-2000

C:Accession: I40276

R:Fukunaga, M.; Hamase, A.

J. Clin. Microbiol. 33, 2415-2420, 1995

A:Title: Outer surface protein C gene sequence analysis of Borrelia burgdorferi sensu lato

A:Reference number: I40269; MUID:96025162

A:Accession: I40276

A:Status: preliminary; translated from GB/EMBL/DBDJ

A:Molecule type: DNA

A:Residues: 1-207 <RES>

A:Cross-references: GB:D49500; NID:G707095; PIDN:BA08460.1; PID:G769687

C:Superfamily: Lyme disease spirochete surface protein C

Query Match
Best Local Similarity 100.0%; Score 52; DB 2; Length 207;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 PVVAESPCKP 10
|||||
Db 198 PVVAESPCKP 207

RESULT 11
S69919

outer surface protein C precursor - Borrelia garinii (strain PTrob)
C:Species: Borrelia garinii
A:Variety: strain PTrob
C:Date: 06-Dec-1996 #sequence_revision 14-Feb-1997 #text_change 26-May-2000
R:Jauris-Heipke, S.; Liegl, G.; Preac-Mursic, V.; Roessler, D.; Schwab, E.; Soutsche
J. Clin. Microbiol. 33, 1860-1866, 1995
A:Title: Molecular analysis of genes encoding outer surface protein C (OspC) of Borrelia burgdorferi
A:Reference number: I40047; MUID:95395018
A:Accession: S69919
A:Status: nucleic acid sequence not shown; translation not shown
A:Molecule type: DNA
A:Residues: 1-207 <JAU>
A:Cross-references: EMBL:X83554; NID:9872027; PIDN:CA58544.1; PID:9872028
A:Experimental source: strain PTrob
A:Note: the nucleotide sequence was submitted to the EMBL Data Library, January 1995
C:Genetics:
A:Gene: ospC
C:Superfamily: Lyme disease spirochete surface protein C
F:1-18/Domain: signal sequence #status predicted <SIG>
F:19-20/Product: outer surface protein C #status predicted <MAT>

Query Match
Best Local Similarity 100.0%; Score 52; DB 2; Length 207;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 PVVAESPCKP 10
|||||
Db 198 PVVAESPCKP 207

RESULT 12
S69924

outer surface protein C precursor - Borrelia garinii (strain Tis1)
C:Species: Borrelia garinii
A:Variety: strain Tis1
C:Date: 15-Feb-1997 #sequence_revision 27-Feb-1997 #text_change 26-May-2000
R:Jauris-Heipke, S.; Liegl, G.; Preac-Mursic, V.; Roessler, D.; Schwab, E.; Soutsche
J. Clin. Microbiol. 33, 1860-1866, 1995
A:Title: Molecular analysis of genes encoding outer surface protein C (OspC) of Borrelia burgdorferi
A:Reference number: I40047; MUID:95395018
A:Accession: S69924
A:Status: nucleic acid sequence not shown
A:Molecule type: DNA
A:Residues: 1-207 <JAU>
A:Cross-references: EMBL:X81525

A:Experimental source: strain Tis1
 R:Roessler, D.
 Submitted to the EMBL Data Library, September 1994
 A:Reference number: S72674
 A:Accession: S72674
 A:Molecule type: DNA
 A:Residues: 1-77, 'VE', 80-207 <ROE>
 A:Cross-references: EMBL:X81525; NID:g804962; PIDN:CAA57245.1; PID:g804963
 C:Genetics:
 A:Gene: ospC
 C:Superfamily: Lyme disease spirochete surface protein C

Query Match 100.0%; Score 52; DB 2; Length 207;
 Best Local Similarity 100.0%; Pred. No. 0.039;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 PVVAESPKP 10
 |||||
 Db 198 PVVAESPKP 207

RESULT 13
 S37727
 Outer surface protein C precursor - Lyme disease spirochete
 C:Species: Borrelia burgdorferi (Lyme disease spirochete)
 C:Date: 20-Feb-1995 #sequence_revision 20-Feb-1995 #text_change 26-May-2000
 C:Accession: S37727
 R:Jauris-Helpe, S.; Fuchs, R.; Motz, M.; Preac-Mursic, V.; Schwab, E.; Soutschek, E.; M.
 Med. Microbiol. Immunol. 182, 37-50, 1993
 A:Title: Genetic heterogeneity of the genes coding for the outer surface protein C (OspC)
 A:Reference number: S37726; MUID:93268136
 A:Accession: S37727
 A>Status: preliminary
 A:Molecule type: DNA
 A:Residues: 1-207 <JAU>
 A:Cross-references: EMBL:X69595; NID:g311393; PIDN:CAA49305.1; PID:g311394
 C:Superfamily: Lyme disease spirochete surface protein C.

Query Match 100.0%; Score 52; DB 2; Length 207;
 Best Local Similarity 100.0%; Pred. No. 0.039;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 PVVAESPKP 10
 |||||
 Db 198 PVVAESPKP 207

RESULT 14
 I40273
 Outer surface protein C precursor - Borrelia afzelii
 C:Species: Borrelia afzelii
 C:Date: 04-Sep-1997 #sequence_revision 04-Sep-1997 #text_change 26-May-2000
 C:Accession: I40273
 R:Fukunaga, M.; Hamase, A.
 J. Clin. Microbiol. 33, 2415-2420, 1995
 A:Title: Outer surface protein C gene sequence analysis of Borrelia burgdorferi sensu la
 A:Reference number: I40269; MUID:96025162
 A:Accession: I40273
 A>Status: preliminary; translated from GB/EMBL/DBJ
 A:Molecule type: DNA
 A:Residues: 1-209 <RES>
 A:Cross-references: GB:D49379; NID:g1041107; PIDN:BA08377.1; PID:g1041108
 C:Superfamily: Lyme disease spirochete surface protein C

Query Match 100.0%; Score 52; DB 2; Length 209;
 Best Local Similarity 100.0%; Pred. No. 0.039;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 PVVAESPKP 10
 |||||

Db 200 PVVAESPKP 209

RESULT 15
 I40270
 Outer surface protein C precursor - Borrelia garinii
 C:Species: Borrelia garinii
 C:Date: 04-Sep-1997 #sequence_revision 04-Sep-1997 #text_change 26-May-2000
 C:Accession: I40270
 R:Fukunaga, M.; Hamase, A.
 J. Clin. Microbiol. 33, 2415-2420, 1995
 A:Title: Outer surface protein C gene sequence analysis of Borrelia burgdorferi ser
 A:Reference number: I40269; MUID:96025162
 A:Accession: I40270
 A>Status: preliminary; translated from GB/EMBL/DBJ
 A:Molecule type: DNA
 A:Residues: 1-209 <RES>
 A:Cross-references: GB:D49498; NID:g707093; PIDN:BA08458.1; PID:g769685
 C:Superfamily: Lyme disease spirochete surface protein C

Query Match 100.0%; Score 52; DB 2; Length 209;
 Best Local Similarity 100.0%; Pred. No. 0.039;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 PVVAESPKP 10
 |||||
 Db 200 PVVAESPKP 209

Search completed: October 12, 2002, 20:45:30
 Job time: 13040 sec

GenCore version 4.5
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OM protein - protein search, using sw model

Run on: October 12, 2002, 20:42:41 ; Search time 35.71 Seconds
(without alignments)
10.843 Million cell updates/sec

Title: US-09-408-578A-1
Perfect score: 52
Sequence: 1 PVAESPCKP 10

Scoring table:
BIOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 105224 seqs, 38719550 residues
Total number of hits satisfying chosen parameters: 105224

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : SwissProt_40:*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	52	100.0	210	1	OSCL_BORBU
2	52	100.0	212	1	OSCL_BORBU
3	37	71.2	205	1	CIP1_RHINE
4	37	71.2	302	1	RP54_AZOVI
5	37	71.2	865	1	CPN_DROME
6	36	69.2	122	1	YRBI_SYNP6
7	36	69.2	382	1	YJOT_YEAST
8	35	69.2	2150	1	SDC3_GABEL
9	35	67.3	297	1	HMXI_BOVIN
10	35	67.3	297	1	HMXI_BOVIN
11	35	67.3	299	1	ODC_HUMAN
12	35	67.3	1388	1	HRP3_SCHPO
13	34	65.4	102	1	VE7_PAPVD
14	34	65.4	387	1	PAB_PEPMA
15	34	65.4	430	1	SYTB_MOUSE
16	34	65.4	430	1	SYTB_MOUSE
17	34	65.4	430	1	SYTB_MOUSE
18	34	65.4	482	1	SYTB_MOUSE
19	34	65.4	486	1	PRLI_ARATH
20	34	65.4	489	1	FLIF_AOUAE
21	34	65.4	528	1	ASMA_YEAST
22	34	65.4	599	1	TRM2_ARATH
23	34	65.4	627	1	DTNB_HUMAN
24	34	65.4	700	1	DTNB_MOUSE
25	34	65.4	1164	1	BAG_STRAG
26	34	65.4	3707	1	PGBM_MOUSE
27	33.5	64.4	226	1	HIC1_XENLA
28	33.5	64.4	220	1	HIC2_XENLA
29	33	63.5	209	1	HIA_XENLA
30	33	63.5	211	1	ROPB_RHILV
31	33	63.5	219	1	H1B_XENLA
32	33	63.5	288	1	HMX3_CHICK
33	33	63.5	333	1	GALF_STMMU

34	33	63.5	372	1	TOLA_HAEIN
35	33	63.5	379	1	PSPB_DICDI
36	33	63.5	388	1	MPRI_YEAST
37	33	63.5	416	1	TRAI_HUMAN
38	33	63.5	472	1	SLX4_DROME
39	33	63.5	492	1	TM62_HUMAN
40	33	63.5	552	1	AAK2_HUMAN
41	33	63.5	552	1	AAK2_RAT
42	33	63.5	587	1	NDC2_RAT
43	33	63.5	616	1	PLRL_RABIT
44	33	63.5	660	1	TPPB_MERTUA
45	33	63.5	885	1	ASE1_YEAST

ALIGNMENTS

RESULT	ID	OSCL_BORBU	STANDARD	PRT	210 AA.
AC	007337				
DT	15-DEC-1998	(Rel. 37, Created)			
DT	15-DEC-1998	(Rel. 37, Last sequence update)			
DT	16-OCT-2001	(Rel. 40, Last annotation update)			
DE	Outer surface protein C precursor (PC).				
GN	OSPC OR BB819.				
OS	Borrelia burgdorferi (Lyme disease spirochete).				
OG	Plasmid lp54.				
OC	Bacteria; Spirochaetales; Spirochaetaceae; Borrelia.				
OX	NCHI_TaxID=139;				
RN	[1]				
RP	SEQUENCE FROM N.A.				
RC	STRAIN-ATCC 35210 / B31;				
RX	MEDLINE-93268136; PubMed-8098841;				
RA	Jauris-Helpke S., Fuchs R., Motz M., Preac-Mursic V., Schwab E.,				
RA	Will G., Wilske B.;				
RT	"Genetic heterogeneity of the genes coding for the outer surface				
RT	protein C (OspC) and the flagellin of Borrelia burgdorferi.";				
RL	Med. Microbiol. Immunol. 182:37-50(1993).				
RN	[2]				
RP	SEQUENCE FROM N.A.				
RC	STRAIN-ATCC 35210 / B31;				
RX	MEDLINE-93239332; PubMed-8478108;				
RA	Wilske B., Preac-Mursic V., Jauris S., Pradel I., Soutschek E.,				
RA	Schwab E., Wanner G.;				
RT	"Immunological and molecular polymorphisms of OspC, an immunodominant				
RT	major outer surface protein of Borrelia burgdorferi.";				
RL	Infect. Immun. 61:2182-2191(1993).				
RN	[3]				
RP	SEQUENCE FROM N.A.				
RC	STRAIN-ATCC 35210 / B31;				
RX	MEDLINE-94041630; PubMed-8225587;				
RA	Padula S.J., Samperi A., Dias F., Szczepanski A., Ryan R.W.;				
RT	"Molecular characterization and expression of p23 (OspC) from a North				
RT	American strain of Borrelia burgdorferi.";				
RL	Infect. Immun. 61:5097-5105(1993).				
RN	[4]				
RP	SEQUENCE FROM N.A.				
RC	STRAIN-ATCC 35210 / B31;				
RX	MEDLINE-96025162; PubMed-7494039;				
RA	Fukunaga M., Hamase A.;				
RT	"Outer surface protein C gene sequence analysis of Borrelia				
RT	burgdorferi sensu lato isolates from Japan.";				
RL	J. Clin. Microbiol. 33:2415-2420(1995).				
RN	[5]				
RP	SEQUENCE FROM N.A.				
RC	STRAIN-ATCC 35210 / B31;				
RX	MEDLINE-98065943; PubMed-9403685;				
RA	Fraser C.M., Casjens S., Huang W.M., Sutton G.G., Clayton R.A.,				
RA	Lachgira R., White O., Ketchum K.A., Dodson R., Hickey E.K., Gwinn M.,				
RA	Dougherty B., Tomb J.F., Fleischmann R.D., Richardson D.,				
RA	Peterson J., Kertlavage A.R., Quackenbush J., Salzberg S., Hanson M.,				
RA	van Vugt R., Palmer N., Adams M.D., Gocayne J.D., Weidman J.,				

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CC -----
DR EMBL; AL591793; CAC47803.1; -
DR InterPro; IPR001907; CLP_protease.
DR Pfam; PF00574; CLP_protease.1.
DR PRINTS; PR00127; CLPPROTEASEP.1.
DR PROSITE; PS00382; CLP_PROTEASE_HIS.1.
DR PROSITE; PS00381; CLP_PROTEASE_SER; FALSE_NEG.
KW Hydrolyase; Serine protease; Complete proteome.
FT ACT_SITE 102 102
FT ACT_SITE 127 127 BY SIMILARITY
SQ SEQUENCE 205 AA; 23179 MW; 7E2D00CAFE15B79 CRC64;

Query Match 71.2%; Score 37; DB 1; Length 205;
Best Local Similarity 100.0%; Pred. No. 12;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

CY 4 AESPKKP 10
DB 57 AESPKKP 63
ID P08623; STANDARD; PRT; 502 AA.
RESULT 4
RP04_AZOVI
AC P08623;
DT 01-AUG-1988 (Rel. 08, Created)
DT 01-AUG-1988 (Rel. 08, Last sequence update)
DT 01-FEB-1994 (Rel. 28, Last annotation update)
DE RNA polymerase sigma-54 factor.
GN RPN OR NTRA.
OS Azobacter vinelandii.
OC Bacteria; Proteobacteria; gamma subdivision; Pseudomonadaceae;
OC Azobacter.
ON NCBI_TaxID=354;
RX [1]
RP SEQUENCE FROM N.A.
RC STRAIN-UM;
RC MEDLINE=8142550; PubMed=3481423;
RA Merrick M.J., Gibbins J., Toukadarian A.;
RT "The nucleotide sequence of the sigma factor gene ntra (rpn) of
RT Azobacter vinelandii: analysis of conserved sequences in Ntra
RT proteins.";
RT Mol. Genet. 210:323-330(1987).
CC -1- FUNCTION: THE SIGMA FACTOR IS AN INITIATION FACTOR THAT PROMOTES
CC ATTACHMENT OF THE RNA POLYMERASE TO SPECIFIC INITIATION SITES AND
CC THEN IS RELEASED.
CC -1- FUNCTION: THIS SIGMA FACTOR IS RESPONSIBLE FOR THE EXPRESSION OF
CC THE NITROGEN FIXATION PROTEINS. THE OPEN COMPLEX (SIGMA-54 AND
CC CORE RNA POLYMERASE) SERVES AS THE RECEPTOR FOR RECEIPT OF THE
CC MELTING SIGNAL FROM THE REMOTELY BOUND ACTIVATOR PROTEIN NIFA FOR
CC THE EXPRESSION OF THE NITROGEN FIXATION PROTEINS.
CC -1- SIMILARITY: BELONGS TO THE SIGMA-54 FACTOR FAMILY.
CC -----
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CC -----
DR EMBL; X05888; CAA29314.1; -
DR PIR; S00720; S00720.
DR InterPro; IPR000394; Sigma54_factor.

DR Pfam; PF00309; Sigma54_factors.1.
DR PRINTS; PR00045; SIGMA54FCT.
DR PROSITE; PS00717; SIGMA54.1; 1.
DR PROSITE; PS00718; SIGMA54.2; 1.
DR PROSITE; PS50044; SIGMA54.3; 1.
KW Transcription regulation; Sigma factor; DNA-directed RNA polymerase;
KW DNA-binding; Nitrogen fixation.
FT DOMAIN 11 38
FT DOMAIN 19 40 GLN-RICH.
FT DOMAIN 186 207 LEUCINE-ZIPPER (POTENTIAL).
FT DOMAIN 391 410 LEUCINE-ZIPPER (POTENTIAL).
FT SITE 479 487 H-1-H MOTIF (POTENTIAL).
FT SITE 479 487 RPN BOX.
SQ SEQUENCE 502 AA; 56917 MW; 699A405BE87D4F10 CRC64;

Query Match 71.2%; Score 37; DB 1; Length 502;
Best Local Similarity 66.7%; Pred. No. 29;
Matches 6; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

CY 2 VVAESPKKP 10
DB 453 IAAENPKKP 461
ID CPN_DROME
ID CPN_DROME STANDARD; PRT; 865 AA.
RESULT 5
CPN_DROME
AC 002910;
DT 01-OCT-1993 (Rel. 27, Created)
DT 01-OCT-1993 (Rel. 27, Last sequence update)
DT 01-FEB-1994 (Rel. 28, Last annotation update)
DE Calphoton.
GN CPN OR CAP.
OS Drosophila melanogaster (Fruit fly).
OC Eukaryota; Metazoa; Arthropoda; Tracheata; Hexapoda; Insecta;
OC Pterygota; Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;
OC Ephydroidea; Drosophilidae; Drosophila.
ON NCBI_TaxID=7227;
RX [1]
RP SEQUENCE FROM N.A.
RC STRAIN-CANTON-S;
RC MEDLINE=93165729; PubMed=8094559;
RA Martin J.H., Benzer S., Rudnicka M., Miller C.A.;
RT "Calphoton: a Drosophila photoreceptor cell calcium-binding protein.";
RT Proc. Natl. Acad. Sci. U.S.A. 90:1531-1535(1993).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN-CANTON-S;
RC MEDLINE=93165730; PubMed=8434015;
RA Ballinger D.G., Xue N., Harshman K.D.;
RT "A drosophila photoreceptor cell-specific protein, calphoton, binds
RT calcium and contains a leucine zipper.";
RT Proc. Natl. Acad. Sci. U.S.A. 90:1536-1540(1993).
CC -1- FUNCTION: MIGHT FUNCTION AS A CALCIUM-SENSITIZING. "SPONGE" TO
CC REGULATE THE AMOUNT OF FREE CYTOPLASMIC CALCIUM. IT BINDS 0.3 MOL
CC OF CA+2 PER MOL OF PROTEIN.
CC -1- SUBUNIT: HOMODIMER (PROBABLE).
CC -1- SUBCELLULAR LOCATION: CYTOPLASMIC; HYPODENSE COMPARTMENT.
CC -1- TISSUE SPECIFICITY: SOMA AND AXONS OF PHOTORECEPTOR CELLS OF
CC COMPOUND EYES AND OCCELLI.
CC -1- DEVELOPMENTAL STAGE: EXPRESSED EARLY IN PHOTORECEPTOR CELL
CC DEVELOPMENT.
CC -----
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CC -----
DR EMBL; L02111; AAA28405.1; -
DR EMBL; L05080; AAA28420.1; -

DR PIR: A47282; A47282.
 DR Flybase: FBgn0010218; Cpn.
 KM Calcium-binding.
 FT CONFLICT 36 36 A -> AVAPAVYA (IN REF. 2).
 FT CONFLICT 43 43 I -> T (IN REF. 2).
 FT CONFLICT 64 64 I -> V (IN REF. 2).
 FT CONFLICT 76 76 T -> A (IN REF. 2).
 FT CONFLICT 100 100 P -> PP (IN REF. 2).
 FT CONFLICT 126 127 VO -> AP (IN REF. 2).
 FT CONFLICT 154 127 I -> V (IN REF. 2).
 FT CONFLICT 160 160 S -> T (IN REF. 2).
 FT CONFLICT 534 534 I -> E (IN REF. 2).
 FT CONFLICT 699 699 I -> T (IN REF. 2).
 FT CONFLICT 703 703 V -> L (IN REF. 2).
 FT CONFLICT 721 721 D -> E (IN REF. 2).
 SQ SEQUENCE 865 AA; 84781 MW; 2110417E0B0E7CFE CRC64;

Query Match
 Best Local Similarity 71.2%; Score 37; DB 1; Length 865;
 Matches 7; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

QY 1 PVVAESPCKP 10
 Db 200 PVVAESPCKP 209

RESULT 6
 YRBL_SYNP6 STANDARD; PRT; 122 AA.
 AC P23655;
 DT 01-NOV-1991 (Rel. 20, Last sequence update)
 DT 16-OCT-2001 (Rel. 40, Last annotation update)
 DE Hypothetical 12.9 kDa protein in RUBISCO 5' region (ORF1).
 OS *Synechococcus* sp. (strain PCC 6301) (*Anacystis nidulans*).
 OC Bacteria; Cyanobacteria; Chroococcales; *Synechococcus*.
 NX NCBI_Taxid=1139;
 RN
 RP SEQUENCE FROM N.A.
 RA Shinozaki K., Sugita M.;
 RT "Genes for the large and small subunits of ribulose-1,5-bisphosphate
 RT carboxylase/oxygenase constitute a single operon in a cyanobacterium
 RT *Anacystis nidulans* 6301.";
 RL Mol. Gen. Genet. 200:27-32(1985).
 CC -1- FUNCTION: MAY BE INVOLVED IN THE FORMATION OF THE CARBOXYISOME, A
 CC POLYMERAL INCLUSION WHERE RUBISCO IS SEQUESTERED.
 CC -1- SIMILARITY: BELONGS TO THE CCM/CCMK/CSO1/PDUA FAMILY.
 CC
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 CC
 CC
 DR EMBL: X03220; CAA26970.1; -;
 DR PIR: S07310; S07310.
 DR InterPro: IPR000249; Bact_microcomp.
 DR Pfam: PF00936; Bact_microcomp; 1.
 DR ProDom: PD003442; Bact_microcomp; 1.
 DR PROSITE: PS01139; BACT_MICROCOMP; 1.
 KW Hypothetical protein.
 SQ SEQUENCE 122 AA; 12855 MW; F54244DE24531504 CRC64;

Query Match
 Best Local Similarity 69.2%; Score 36; DB 1; Length 122;
 Matches 6; Conservative 1; Mismatches 3; Indels 0; Gaps 0;
 QY 1 PVVAESPCKP 10
 Db 200 PVVAESPCKP 209

Db 100 PIAGSPCKP 109

RESULT 7
 YJ07_YEAST STANDARD; PRT; 382 AA.
 ID YJ07_YEAST
 AC P47007;
 DT 01-FEB-1996 (Rel. 33, Created)
 DT 01-FEB-1996 (Rel. 33, Last sequence update)
 DT 01-FEB-1996 (Rel. 33, Last annotation update)
 DE Hypothetical 44.9 kDa protein in INO1-IDS2 intergenic region.
 GN YJL147C OR J0639.
 OS *Saccharomyces cerevisiae* (Baker's yeast).
 OS *Saccharomyces cerevisiae*.
 OC Eukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomycetes;
 OC Saccharomycetales; Saccharomycetaceae; Saccharomycetes.
 NX NCBI_Taxid=4932;
 RN
 RP SEQUENCE FROM N.A.
 RC STRAIN=5288C / FY1679;
 RX MEDLINE=96408771; PubMed=8813765;
 RA Katsoulou C., Tzeremia M., Tavernarakis N., Alexandraki D.;
 RT "Sequence analysis of a 40.7 kb segment from the left arm of yeast
 RT chromosome X reveals 14 known genes and 13 new open reading frames
 RT including homologues of genes clustered on the right arm of
 RT chromosome XI.";
 RL Yeast 12:787-797(1996).
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 CC
 CC
 DR EMBL: Z49422; CAA89442.1; -;
 DR EMBL: X87371; CAA60808.1; -;
 DR SGD: S0003663; YJL147C.
 KW Hypothetical protein.
 SQ SEQUENCE 382 AA; 44862 MW; DDAAAF588AF9A3234 CRC64;

Query Match
 Best Local Similarity 69.2%; Score 36; DB 1; Length 382;
 Matches 6; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 2 VVAESPCKP 10
 Db 313 IMLESPCKP 321

RESULT 8
 SDC3_CAEL STANDARD; PRT; 2150 AA.
 ID SDC3_CAEL
 AC P34706;
 DT 01-FEB-1994 (Rel. 28, Created)
 DT 01-FEB-1994 (Rel. 28, Last sequence update)
 DT 01-MAR-2002 (Rel. 41, Last annotation update)
 DE Zinc finger protein Sdc-3.
 GN SDC-3.
 OS *Caenorhabditis elegans*.
 OC Eukaryota; Metazoa; Nematoda; Chromadorea; Rhabditida; Rhabditidae;
 OC Rhabditidae; Peloderinae; Caenorhabditis.
 NX NCBI_Taxid=6239;
 RN
 RP SEQUENCE FROM N.A.
 RC STRAIN=BRISTOL N2;
 RX MEDLINE=93161411; PubMed=8431944;
 RA Klein R.D., Meyer B.J.;
 RT "Independent domains of the Sdc-3 protein control sex determination
 RT and dosage compensation in *C. elegans*.";
 RL Cell 72:349-364(1993).
 CC -1- FUNCTION: CONTROLS BOTH SEX DETERMINATION AND X CHROMOSOME DOSAGE

```

DR EMBL: D30750; BAA20367.1; -
DR HSSP: P22608; INK3
DR InterPro; IPR001356; Homeobox.
DR Pfam; PF00046; homeobox; 1.
DR PRINTS; PR00024; HOMEBOX.
DR SMART; SM00389; HOX; 1.
DR PROSITE; PS00027; HOMEBOX_1; 1.
DR PROSITE; PS0071; HOMEBOX_2; 1.
DR Homeobox; DNA-binding; Developmental protein; Nuclear protein.
KT DNA_BIND 166 225
SQ SEQUENCE 297 AA; 31111 MW; 4D0C987EE1F6FBEB CRC64;

Query Match 67.3%; Score 35; DB 1; Length 297;
Best Local Similarity 77.8%; Pred. No. 40;
Matches 7; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

OY 2 VYAESPKRP 10
      1 11111111
DB 126 VKAESPEKP 134

RESULT 10
HMX1_HUMAN
ID HMX1_HUMAN STANDARD; PRT; 297 AA.
AC P28360; Q96NY4;
DT 01-DEC-1992 (Rel. 24, Created)
DT 01-DEC-1992 (Rel. 24, Last sequence update)
DT 01-MAR-2002 (Rel. 41, Last annotation update)
DE Homeobox protein MSX-1 (Hox-7).
GN MSX1 OR HOX7.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
OX NCBI_TaxID=9606;

[1]
SEQUENCE FROM N.A.
MEDLINE=93250782; PubMed=1284527;
RA Padanilam B., Stedler S.H., Mills K.A., McLeod L.B., Solursh M.,
RA Lee B.M., Ramirez F., Bucetow K.H., Murray J.C.;
RT "Characterization of the human HOX 7 cDNA and identification of
RT polymorphic markers.";
RL Hum. Mol. Genet. 1:407-410(1992).

[2]
SEQUENCE FROM N.A.
MEDLINE=92128949; PubMed=1685479;
RA Hewitt J.E., Clarke L.E., Iyen A., Williamson R.;
RT "Structure and sequence of the human homeobox gene HOX7.";
RL Genomics 11:670-678(1991).

[3]
SEQUENCE FROM N.A.
MEDLINE=96331281; PubMed=8696335;
RA Vestergaard H., Karimibay N., Guthrie S.W., Seidman J.G., Seidman C.E.;
RT "A human MSX1 homeobox domain missense mutation causes selective tooth
RT agenesis.";
RL Nat. Genet. 13:417-421(1996).

[4]
VARIANT FTA PRO-196.
RX Vestergaard H., Karimibay N., Guthrie S.W., Seidman J.G., Seidman C.E.;
RT "Diagnostic resequencing demonstrates a role for MSX1 in nonsyndromic
RT cleft lip and palate.";
RL Submitted (OCT-2001) to the EMBL/GenBank/DBJ databases.

[5]
VARIANT FTA PRO-196.
RX Vestergaard H., Karimibay N., Guthrie S.W., Seidman J.G., Seidman C.E.;
RT "A human MSX1 homeobox domain missense mutation causes selective tooth
RT agenesis.";
RL Nat. Genet. 13:417-421(1996).

[6]
FUNCTION: PROBABLE MORPHOGENETIC ROLE. MAY PLAY A ROLE IN LIMB-
PATTERN FORMATION.
[7]
SUBCELLULAR LOCATION: Nuclear.
[8]
DISEASE: IMPLICATED IN WOLF-HIRSCHORN SYNDROME (WHS), WHICH IS
CHARACTERIZED BY PROFOUND MENTAL RETARDATION, HEART DEFECTS, AND
FACIAL CLEFTING.
[9]
DISEASE: DEFECTS IN MSX1 ARE IMPLICATED IN FAMILIAL TOOTH AGENESIS
(FTA). AGENESIS OF ONE OR MORE TEETH CONSTITUTES ONE OF THE MOST
COMMON DEVELOPMENTAL ANOMALIES IN MAN. REPORTED INCIDENCES VARY
FROM 1.6% TO 9.6%, EXCLUDING THIRD MOLAR (WISDOM TOOTH) AGENESIS,
WHICH OCCURS IN 70% OF THE POPULATION.

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-1- SIMILARITY: BELONGS TO THE MSH FAMILY OF HOMEBOX PROTEINS.
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CC -----
DR EMBL; M97676; AAA52683.1; -.
DR EMBL; M76732; AAA58665.1; -.
DR EMBL; M76731; AAA58665.1; JOINED.
DR EMBL; AF426432; AA117870.1; -.
DR PIR; A40560; A40560.
DR HSSP; P22808; INK3.
DR TRANSFAC; T02071; -.
DR MIM; 142963; -.
DR MIM; 106600; -.
DR InterPro; IPR001827; Antennapedia.
DR InterPro; IPR001356; Homeobox.
DR Pfam; PF00046; homeobox; 1.
DR PRINTS; PRO0025; ANTENNAPEDIA.
DR PRINTS; PRO0024; HOMEBOX.
DR SMART; SMO0389; HOX; 1.
DR PROSITE; PS00027; HOMEBOX_1; 1.
DR PROSITE; PS50071; HOMEBOX_2; 1.
DR Homeobox; DNA-binding; Developmental protein; Nuclear protein;
DR disease mutation.
FT FT DNA_BIND 166 225 HOMEBOX:
FT FT VARIANT 196 196 R->P (IN FTN).
FT FT CONFLICT 39 39 /FTID=VAR_003754.
FT FT CONFLICT 91 92 A->T (IN REF. 2).
FT FT CONFLICT 93 93 GV->AS (IN REF. 2).
SO SEQUENCE 297 AA; 30998 MW; A19B4F0E39567B41 CRC64;

Query Match 67.3%; Score 35; DB 1; Length 297;
Best Local Similarity 77.8%; Pred. No. 40;
Matches 7; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 2 VVASEPKKP 10
| | | | | | | |
DB 126 VKASEPEK 134

RESULT 11
ODC_HUMAN
AC O9B0T8; STANDARD; PRT; 299 AA.
DT 01-MAR-2002 (Rel. 41, Created)
DT 01-MAR-2002 (Rel. 41, Last sequence update)
DT 01-MAR-2002 (Rel. 41, Last annotation update)
DE Mitochondrial 2-oxodicarboxylate carrier.
GN SLC25A21 OR ODC.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Euthera; Primates; Catarrhini; Homnidae; Homo.
OX NCBI_TaxId=9606;
RN [1]
RP SEQUENCE FROM N.A., CHARACTERIZATION, AND TISSUE SPECIFICITY.
RC TISSUE=Liver;
RA MEDLINE=21269385; PubMed=11083877;
RA Fiermonte G., Dolce V., Palmieri L., Ventura M., Runswick M.J.,
RA Palmieri F., Walker J.E.;
RT "Identification of the human mitochondrial oxodicarboxylate carrier:
RT bacterial expression, reconstruction, functional characterization,
RT tissue distribution and chromosomal location.";
RT J. Biol. Chem. 276:8225-8230(2001).
FT FUNCTION: Transports C5-C7 oxodicarboxylates across the inner
CC membranes of mitochondria. Can transport 2-oxoadipate, 2-
CC oxoglutarate, adipate, glutarate, and to a lesser extent,
CC

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CC plimete, 2-oxopimelate, 2-aminoadipate, oxaloacetate, and
CC citrate.
CC
CC -1- SUBCELLULAR LOCATION: Integral membrane protein. Mitochondrial
CC inner membrane.
CC -1- DOMAIN: COMPOSED OF THREE HOMOLOGOUS DOMAINS.
CC -1- TISSUE SPECIFICITY: Ubiquitous.
CC -1- SIMILARITY: BELONGS TO THE MITOCHONDRIAL CARRIER FAMILY.
CC -----
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CC -----
CC EMBL, AJ278148; CAC27562.1;
CC DR Interpro; IPR001993; Mitoch_carrier.
CC DR Interpro; IPR002067; Mit_carrier.
CC DR Pfam; PF00153; mito_carr; 3.
CC DR PRINTS; PR00926; MITOCARRIER.
CC DR PROSITE; PS00215; MITOCH_CARRIER; 2.
CC KW Mitochondrion; Inner membrane; Repeat; Transmembrane; Transport.
CC FT TRANSMEM 17 37 POTENTIAL.
CC FT TRANSMEM 62 82 POTENTIAL.
CC FT TRANSMEM 100 120 POTENTIAL.
CC FT TRANSMEM 211 231 POTENTIAL.
CC SO SEQUENCE 299 AA; 33303 MW; 69A259400328AE19 CRC64;
CC
CC Query Match 67.3%; Score 35; DB 1; Length 299;
CC Best Local Similarity 55.6%; Pred. No. 40;
CC Matches 5; Conservative 4; Mismatches 0; Indels 0; Gaps 0;
CC
CC QY 1 PVVASEPKK 9
CC Db 79 PILAETPKR 87
CC
CC RESULT 12
CC HRP3_SCHPO
CC ID HRP3_SCHPO STANDARD; PRG: 1368 AA.
CC OL1139;
CC DT 16-OCT-2001 (Rel. 40, Created)
CC DT 16-OCT-2001 (Rel. 40, Last sequence update)
CC DT 16-OCT-2001 (Rel. 40, Last annotation update)
CC DE Chromodomain helicase hrp3.
CC GN HRP3 OR SPAC36.01.
CC OS Schizosaccharomyces pombe (Fission yeast).
CC OC Eukaryota; Fungi; Ascomycota; Schizosaccharomycetes;
CC OC Schizosaccharomycetales; Schizosaccharomycetaceae;
CC OC Schizosaccharomycetes.
CC OX NCBI_TaxID=4896;
CC RN [1]
CC RA SEQUENCE FROM N.A.
CC RC SPRAIN-972;
CC RA Gentles S., Churcher C.M., Barrell B.G., Rajandream M.A., Wood V.;
CC RN submitted (SEP-1997) to the EMBL/GenBank/DBJ databases.
CC [2]
CC RA GENE NAME.
CC RP Bjerling P., Ekwall K.;
CC RA Submitted (MAR-2001) to the SWISS-PROT data bank.
CC CC -1- SUBCELLULAR LOCATION: Nuclear (Potential);
CC CC -1- SIMILARITY: BELONGS TO THE SRF2/RAD54 HELICASE FAMILY.
CC CC -1- SIMILARITY: CONTAINS 2 'CHROMO' DOMAINS.
CC -----
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CC EMBL: Z99167; CAB16277.1;
DR InterPro: IPR000953; Chromo.
DR InterPro: IPR001410; DEAD.
DR InterPro: IPR001650; Helicase_C.
DR Pfam: PF00385; Chromo; 2.
DR Pfam: PF00271; Helicase_C; 1.
DR Pfam: PF00176; SNF2_N; 1.
DR SMART: SM00298; CHROMO; 2.
DR SMART: SM00487; DEXDC; 1.
DR SMART: SM00490; HELIC_C; 1.
DR PROSITE: PS00598; CHROMO_1; 1.
DR PROSITE: PS0013; CHROMO_2; 2.
KM ATP-binding; Helicase; DNA-binding; Nuclear protein; Repeat.
FT DOMAIN 169 175 POLY-GLU.
FT DOMAIN 191 260 CHROMO 1.
FT DOMAIN 288 349 CHROMO 2.
FT NP_BIND 400 407 ATP (POTENTIAL).
FT SITE 508 511 DEAD BOX.
FT DOMAIN 947 950 POLY-GLU.
SQ SEQUENCE 1388 AA; 159377 MW; F7B431084BD29F8D CRC64;

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Query Match
Best Local Similarity 67.3%; Score 35; DB 1; Length 1388;
Matches 6; Conservative 2; Mismatches 2; Indels 0; Gaps 0;
QY 1 PYVAESPKKP 10
Db 1251 PAISESRKRP 1260

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RESULT 13
VE7_PAPVD STANDARD; PRT; 102 AA.
ID P03131;
AC 21-JUL-1986 (Rel. 01, Created)
DT 01-FEB-1996 (Rel. 33, Last sequence update)
DT 01-MAR-2002 (Rel. 41, Last annotation update)
DE E7 protein.
GN E7.
OS Deer papillomavirus.
OC Viruses; dsDNA viruses, no RNA stage; Papillomaviridae;
OC Papillomavirus.
OX NCBI_TaxID=10564;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=85293253; PubMed=2993669;
RA Groff D.E., Lancaster W.D.;
RT "Molecular cloning and nucleotide sequence of deer papillomavirus.";
RL J. Virol. 56:85-91(1985).
CC -1- FUNCTION: E7 PROTEIN HAS BOTH TRANSFORMING AND TRANS-ACTIVATING
ACTIVITIES.
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CC -----
CC EMBL: M11910; AAA6842.1;
CC PIR: A03693; W7MLDP.
CC InterPro: IPR000148; Papyl_E7.
CC Pfam: PF00527; E7; 1.
CC Early protein; Transcription regulation; Oncogene;
CC DNA-binding; Trans-acting factor.
FT DOMAIN 61 64 C-X-X-C MOTIF 1.
FT DOMAIN 94 97 C-X-X-C MOTIF 2.
SQ SEQUENCE 102 AA; 11166 MW; CEAB37C8955FC03 CRC64;

```

```

Query Match
Best Local Similarity 65.4%; Score 34; DB 1; Length 102;
Matches 6; Conservative 1; Mismatches 3; Indels 0; Gaps 0;
QY 1 PYVAESPKKP 10
Db 37-PVYVDKPKP 46

```

```

RESULT 14
PAB_PEPMA STANDARD; PRT; 387 AA.
ID PAB_PEPMA
AC Q51911;
DT 15-JUL-1998 (Rel. 36, Created)
DT 15-JUL-1998 (Rel. 36, Last sequence update)
DE Peptostreptococcal albumin-binding protein precursor.
GN PAB.
OS Peptostreptococcus magnus.
OC Bacteria; Firmicutes; Bacillus/Clostridium group; Clostridiaceae;
OC Finegoldia.
OX NCBI_TaxID=1260;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=ALB8;
RX MEDLINE=94216330; PubMed=8163519;
RA de Chateau M., Björck L.;
RT "Protein PAB, a mosaic albumin-binding bacterial protein representing
RT the first contemporary example of module shuffling."
RL J. Biol. Chem. 269:12147-12151(1994).
RN [2]
RP STRUCTURE BY NMR OF 213-265, AND REVISION TO 244.
RC STRAIN=ALB8;
RX MEDLINE=97240805; PubMed=9086265;
RA Johansson M.O., de Chateau M., Wikström M., Forsen S., Drakenberg T.,
RA Björck L.;
RT "Solution structure of the albumin-binding GA module: a versatile
RT bacterial protein domain."
RL J. Mol. Biol. 266:859-865(1997).
CC -1- FUNCTION: BINDS SERUM ALBUMIN.
CC -----
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CC or send an email to license@isb-sib.ch).
CC -----
CC EMBL: X77864; CA54857.1;
CC PDB: 1GAB; 07-JUL-97.
CC PDB: 1PRB; 23-JUL-97.
CC InterPro: IPR002988; GA.
CC InterPro: IPR001899; Gram_pos_anchor.
CC Pfam: PF01468; GA; 2.
CC Pfam: PF00746; Gram_pos_anchor; 1.
KW signal; 3D-structure.
FT SIGNAL 1 26
FT CHAIN 27 387 PEPTOSTREPTOCOCCAL ALBUMIN-BINDING
FT PROTEIN.
FT DOMAIN 213 265 GA MODULE.
SQ SEQUENCE 387 AA; 43057 MW; 3D5135C4C3A3BD8F2 CRC64;

```

```

Query Match
Best Local Similarity 65.4%; Score 34; DB 1; Length 387;
Matches 6; Conservative 1; Mismatches 0; Indels 0; Gaps 0;
QY 4 AESPKKP 10
Db 194-AETPKKP 200

```

```
RESULT 15
SYTB_MOUSE
ID SYTB_MOUSE STANDARD; PRT; 430 AA.
AC Q9RON3;
DT 16-OCT-2001 (Rel. 40, Created)
DT 16-OCT-2001 (Rel. 40, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE Synaptotagmin XI (SYTXI).
GN SYT11.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=ICR; TISSUE=Cerebellum;
RX MEDLINE=20002669; PubMed=10531343;
RA Fukuda M., Kanno E., Mikoshiba K.;
RT "Conserved N-terminal cysteine motif is essential for homo- and
RT heterodimer formation of synaptotagmins III, V, VI, and X.";
RL J. Biol. Chem. 274:31421-31427(1999).
CC -1- FUNCTION: MAY BE INVOLVED IN CA2+-DEPENDENT EXOCYTOSIS OF
CC SECRETORY VESICLES THROUGH CA2+ AND PHOSPHOLIPID BINDING TO THE C2
CC DOMAIN OR MAY SERVE AS CA2+ SENSORS IN THE PROCESS OF VESICULAR
CC TRAFFICKING AND EXOCYTOSIS (By similarity).
CC -1- SUBUNIT: Homodimer. Can also forms heterodimer (By similarity).
CC -1- SUBCELLULAR LOCATION: INTEGRAL MEMBRANE PROTEIN. SYNAPTIC
CC VESICLES (By similarity).
CC -1- SIMILARITY: CONTAINS 2 C2 DOMAINS.
CC -1- SIMILARITY: BELONGS TO THE SYNAPTOTAGMIN FAMILY.
CC -----
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CC -----
DR EMBL; AB026808; BAA85780.1; -.
DR HSSP; P21707; IRSY.
DR MGD; MGI:1859547; SYT11.
DR InterPro; IPR000008; C2.
DR InterPro; IPR002149; LRI.
DR InterPro; IPR001565; Synaptotagmin.
DR Pfam; PF00168; C2; 2.
DR PRINTS; PR00360; C2DOMAIN.
DR PRINTS; PR00399; SYNAPTOTAGMIN.
DR SMART; SM00239; C2; 2.
DR PROSITE; PS00499; C2_DOMAIN_1; FALSE_NEG.
DR PROSITE; PS50004; C2_DOMAIN_2; 2.
KW Transmembrane; Repeat; Synapse.
FT DOMAIN 1 15 VESICULAR (POTENTIAL).
FT TRANSMEM 16 36 POTENTIAL.
FT DOMAIN 37 430 CYTOPLASMIC (POTENTIAL).
FT DOMAIN 173 261 C2 DOMAIN 1.
FT DOMAIN 303 396 C2 DOMAIN 2.
SQ SEQUENCE 430 AA; 48359 MW; 25E7CDFC4B4BE036 CRC64;
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Query Match 65.4%; Score 34; DB 1; Length 430;
Best Local Similarity 75.0%; Pred. No. 86;
Matches 6; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 3 VAESPGRP 10
| | | | |
Db 413 VCESPRKP 420

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OM protein - protein search, using sw model

Run on: October 12, 2002, 17:27:46 ; Search time 87.59 Seconds

(without alignments)
19.751 Million cell updates/sec

Title: US-09-408-578A-1

Perfect score: 52

Sequence: 1 PVVAESPKRP 10

Scoring table: BIOSUM62

Searched: Gapop 10.0, Gapext 0.5

Total number of hits satisfying chosen parameters: 562222

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :
1: SP archaea:*
2: SP bacteria:*
3: SP fungi:*
4: SP human:*
5: SP invertebrate:*
6: SP mammal:*
7: SP mhc:*
8: SP organelle:*
9: SP phage:*
10: SP plant:*
11: SP rodent:*
12: SP virus:*
13: SP vertebrate:*
14: SP unclassified:*
15: SP virus:*
16: SP bacteriophage:*
17: SP archaea:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	52	100.0	190	2	P94244
2	52	100.0	190	2	P70819
3	52	100.0	191	2	P94245
4	52	100.0	191	2	O9S3P0
5	52	100.0	191	2	P70818
6	52	100.0	192	2	O9S3P3
7	52	100.0	192	2	O9S3P2
8	52	100.0	193	2	P94226
9	52	100.0	193	2	P94233
10	52	100.0	194	2	P94237
11	52	100.0	194	2	O9S3P4
12	52	100.0	194	2	P94229
13	52	100.0	194	2	O45030
14	52	100.0	207	2	O45187
15	52	100.0	207	2	O45177
16	52	100.0	207	2	O45177

17	52	100.0	207	2	O49581	O49581 borrelia ga
18	52	100.0	207	2	O45175	O45175 borrelia ga
19	52	100.0	207	2	O07336	O07336 borrelia bu
20	52	100.0	209	2	O44883	O44883 borrelia bu
21	52	100.0	209	2	O45179	O45179 borrelia ga
22	52	100.0	209	2	O9K1K3	O9K1K3 borrelia ga
23	52	100.0	209	2	O44671	O44671 borrelia af
24	52	100.0	209	2	O49579	O49579 borrelia ga
25	52	100.0	209	2	O49583	O49583 borrelia ga
26	52	100.0	209	2	O49584	O49584 borrelia ga
27	52	100.0	209	2	P70891	P70891 borrelia ga
28	52	100.0	210	2	O57279	O57279 borrelia ga
29	52	100.0	210	2	O57359	O57359 borrelia ga
30	52	100.0	210	2	O9K1M6	O9K1M6 borrelia bu
31	52	100.0	210	2	O45176	O45176 borrelia ga
32	52	100.0	210	2	O45178	O45178 borrelia ga
33	52	100.0	210	2	O49582	O49582 borrelia ga
34	52	100.0	210	2	P70893	P70893 borrelia ga
35	52	100.0	210	2	O44719	O44719 borrelia bu
36	52	100.0	211	2	O44978	O44978 borrelia bu
37	52	100.0	211	2	O57262	O57262 borrelia af
38	52	100.0	211	2	O49576	O49576 borrelia af
39	52	100.0	211	2	O49577	O49577 borrelia af
40	52	100.0	211	2	O49576	O49576 borrelia bu
41	52	100.0	211	2	O44720	O44720 borrelia bu
42	52	100.0	211	2	O44977	O44977 borrelia bu
43	52	100.0	212	2	O926C7	O926C7 borrelia va
44	52	100.0	212	2	O9K1M5	O9K1M5 borrelia af
45	52	100.0	212	2	O9K1M4	O9K1M4 borrelia af

ALIGNMENTS

RESULT 1
P94244 PRELIMINARY; PRT; 190 AA.
AC P94244;
DT 01-MAY-1997 (TREMBLrel. 03, Created)
DT 01-MAY-1997 (TREMBLrel. 03, Last sequence update)
DT 01-JUN-2001 (TREMBLrel. 17, Last annotation update)
DE OUTER SURFACE PROTEIN C (FRAGMENT).
GN OSpC.
OS Borrelia burgdorferi (Lyme disease spirochete).
OC Bacteria; Spirochaetales; Spirochaetaceae; Borrelia.
OX NCBI_TaxID-139;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN-VSDA;
RX MEDLINE-96296448; PubMed-8709845;
RA Livey, I., Gibbs C.P., Schuster R., Dorner F.;
RT "Evidence for lateral transfer and recombination in OSpC variation in
RT Lyme disease Borrelia".
RL MOL. Microbiol. 18:257-269(1995).
DR EMBL; U42870; AAB37013.1;
DR InterPro; IPR001800; Lipoprotein_6.
DR Pfam; PF01441; Lipoprotein_6; 1.
DR Prodom; PD001149; Lipoprotein_6; 1.
FT NON_TER 1
FT NON_TER 190
SO SEQUENCE 190 AA; 19904 MW; 9A0C47B29B73ADB8 CRC64;

Query Match 100.0%; Score 52; DB 2; Length 190;
Best Local Similarity 100.0%; Pred. No. 0.076;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 1 PVVAESPKRP 10
Db 181 PVVAESPKRP 190

RESULT 2

P70819 PRELIMINARY: PRT: 190 AA.
 AC P70819: (TREMblrel. 02, Created)
 DT 01-FEB-1997 (TREMblrel. 02, Last sequence update)
 DT 01-FEB-1997 (TREMblrel. 03, Last sequence update)
 DT 01-DEC-2001 (TREMblrel. 19, Last annotation update)
 DE OUTER SURFACE PROTEIN (FRAGMENT).
 GN OSC.
 OS Borrelia burgdorferi (Lyme disease spirochete).
 OC Bacteria; Spirochaetales; Spirochaetaceae; Borrelia.
 OX NCBI_TaxID=139;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=2-1498 SON 188;
 RA Probert W.S., Crawford M.R., Cadiz R.B., Lefebvre R.B.;
 RT "Immunization with OspA, but not ospC, provides protection of mice
 challenged with North American isolates of Borrelia burgdorferi.";
 RL Submitted (SEP-1996) to the EMBL/GenBank/DBJ databases.
 DR EMBL: L81130; AAB06570.1; -;
 DR InterPro: IPR001800; Lipoprotein_6.
 DR Pfam: PF01441; Lipoprotein_6; 1.
 DR Prodom: PD001149; Lipoprotein_6; 1.
 FT NON_TER 1
 SQ SEQUENCE 190 AA; 19695 MW; FBB92D4494F2D49A CRC64;

Query Match 100.0%; Score 52; DB 2; Length 190;
 Best Local Similarity 100.0%; Pred. No. 0.076;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 PVVAESPCKP 10
 DB 181 PVVAESPCKP 190

RESULT 3
 P94245 PRELIMINARY: PRT: 191 AA.
 AC P94245: (TREMblrel. 03, Created)
 DT 01-MAY-1997 (TREMblrel. 03, Last sequence update)
 DT 01-MAY-1997 (TREMblrel. 03, Last sequence update)
 DT 01-JUN-2001 (TREMblrel. 17, Last annotation update)
 DE OUTER SURFACE PROTEIN C (FRAGMENT).
 GN OSC.
 OS Borrelia burgdorferi (Lyme disease spirochete).
 OC Bacteria; Spirochaetales; Spirochaetaceae; Borrelia.
 OX NCBI_TaxID=139;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=VS461;
 RA MEDLINE=96296448; PubMed=8709845;
 RA Lyley I., Gibbs C.P., Schuster R., Dorner F.;
 RT "Evidence for lateral transfer and recombination in Ospa variation in
 Lyme disease Borrelia.";
 RL Mol. Microbiol. 18:257-269(1995).
 DR EMBL: L42871; AAB37014.1; -;
 DR InterPro: IPR001800; Lipoprotein_6.
 DR Pfam: PF01441; Lipoprotein_6; 1.
 DR Prodom: PD001149; Lipoprotein_6; 1.
 FT NON_TER 1
 FT NON_TER 191
 SQ SEQUENCE 191 AA; 19923 MW; 2C2F3BD40714EAA8 CRC64;

Query Match 100.0%; Score 52; DB 2; Length 191;
 Best Local Similarity 100.0%; Pred. No. 0.076;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 PVVAESPCKP 10
 DB 182 PVVAESPCKP 191

RESULT 4
 Q9S3P0 PRELIMINARY: PRT: 191 AA.
 AC Q9S3P0: (TREMblrel. 13, Created)
 DT 01-MAY-2000 (TREMblrel. 13, Last sequence update)
 DT 01-MAY-2000 (TREMblrel. 13, Last sequence update)
 DT 01-JUN-2001 (TREMblrel. 17, Last annotation update)
 DE OUTER SURFACE PROTEIN C (FRAGMENT).
 GN OSC.
 OS Borrelia burgdorferi (Lyme disease spirochete).
 OC Bacteria; Spirochaetales; Spirochaetaceae; Borrelia.
 OX NCBI_TaxID=139;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=27579;
 RA Lyley I., Gibbs C.P., Schuster R., Dorner F.;
 RT "Evidence for lateral transfer and recombination in Ospa variation in
 Lyme disease Borrelia.";
 RL Mol. Microbiol. 18:257-269(1995).
 DR EMBL: L42896; AAB37004.1; -;
 DR InterPro: IPR001800; Lipoprotein_6.
 DR Pfam: PF01441; Lipoprotein_6; 1.
 DR Prodom: PD001149; Lipoprotein_6; 1.
 FT NON_TER 1
 FT NON_TER 191
 SQ SEQUENCE 191 AA; 19826 MW; DBA79667814F290A CRC64;

Query Match 100.0%; Score 52; DB 2; Length 191;
 Best Local Similarity 100.0%; Pred. No. 0.076;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 PVVAESPCKP 10
 DB 182 PVVAESPCKP 191

RESULT 5
 P70818 PRELIMINARY: PRT: 191 AA.
 AC P70818: (TREMblrel. 02, Created)
 DT 01-FEB-1997 (TREMblrel. 02, Last sequence update)
 DT 01-FEB-1997 (TREMblrel. 02, Last sequence update)
 DT 01-DEC-2001 (TREMblrel. 19, Last annotation update)
 DE OUTER SURFACE PROTEIN (FRAGMENT).
 GN OSC.
 OS Borrelia burgdorferi (Lyme disease spirochete).
 OC Bacteria; Spirochaetales; Spirochaetaceae; Borrelia.
 OX NCBI_TaxID=139;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC STRAIN=2-1498 CA4;
 RA Probert W.S., Crawford M.R., Cadiz R.B., Lefebvre R.B.;
 RT "Immunization with OspA, but not ospC, provides protection of mice
 challenged with North American isolates of Borrelia burgdorferi.";
 RL Submitted (SEP-1996) to the EMBL/GenBank/DBJ databases.
 DR EMBL: L81131; AAB06569.1; -;
 DR InterPro: IPR001800; Lipoprotein_6.
 DR Pfam: PF01441; Lipoprotein_6; 1.
 DR Prodom: PD001149; Lipoprotein_6; 1.
 FT NON_TER 1
 FT NON_TER 191
 SQ SEQUENCE 191 AA; 20126 MW; D2B9B1C82B4DC3C0 CRC64;

Query Match 100.0%; Score 52; DB 2; Length 191;
 Best Local Similarity 100.0%; Pred. No. 0.076;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 PVVAESPCKP 10
 DB 182 PVVAESPCKP 191

RESULT 6
ID Q9S3P3 PRELIMINARY; PRT; 192 AA.
AC Q9S3P3;
DT 01-MAY-2000 (TREMBLrel. 13, Created)
DT 01-MAY-2000 (TREMBLrel. 13, Last sequence update)
DT 01-JUN-2001 (TREMBLrel. 17, Last annotation update)
DE OUTER SURFACE PROTEIN C (FRAGMENT).
GN OSC.
OS Borrelia burgdorferi (Lyme disease spirochete).
OC Bacteria; Spirochaetales; Spirochaetaceae; Borrelia.
OX NCBI_TaxID=139;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=J1;
RX MEDLINE=96296448; PubMed=8709845;
RA Livey I., Gibbs C.P., Schuster R., Dornier F.;
RT "Evidence for lateral transfer and recombination in OspC variation in
Lyme disease Borrelia";
RL Mol. Microbiol. 18:257-269(1995).
DR EMBL; L42887; AAB36992.1;
DR InterPro: IPR001800; Lipoprotein_6.
DR Pfam: PF01441; Lipoprotein_6; 1.
DR ProDom: PD001149; Lipoprotein_6; 1.
FT NON_TER 1
FT 1
SQ SEQUENCE 192 AA; 20287 MW; 11846F7AC84C7E3D CRC64;

Query Match 100.0%; Score 52; DB 2; Length 192;
Best Local Similarity 100.0%; Pred. No. 0.077;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 PVVAESPKKP 10
Db 183 PVVAESPKKP 192
|||||

RESULT 7
ID Q9S3P2 PRELIMINARY; PRT; 192 AA.
AC Q9S3P2;
DT 01-MAY-2000 (TREMBLrel. 13, Created)
DT 01-MAY-2000 (TREMBLrel. 13, Last sequence update)
DT 01-JUN-2001 (TREMBLrel. 17, Last annotation update)
DE OUTER SURFACE PROTEIN C (FRAGMENT).
GN OSC.
OS Borrelia burgdorferi (Lyme disease spirochete).
OC Bacteria; Spirochaetales; Spirochaetaceae; Borrelia.
OX NCBI_TaxID=139;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=297;
RX MEDLINE=96296448; PubMed=8709845;
RA Livey I., Gibbs C.P., Schuster R., Dornier F.;
RT "Evidence for lateral transfer and recombination in OspC variation in
Lyme disease Borrelia";
RL Mol. Microbiol. 18:257-269(1995).
DR EMBL; L42893; AAB37001.1;
DR InterPro: IPR001800; Lipoprotein_6.
DR Pfam: PF01441; Lipoprotein_6; 1.
DR ProDom: PD001149; Lipoprotein_6; 1.
FT NON_TER 1
FT 1
SQ SEQUENCE 192 AA; 20472 MW; 46AC8F93E4DFED6C CRC64;

Query Match 100.0%; Score 52; DB 2; Length 192;
Best Local Similarity 100.0%; Pred. No. 0.077;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 PVVAESPKKP 10

Db 183 PVVAESPKKP 192
|||||
RESULT 8
ID P94226 PRELIMINARY; PRT; 193 AA.
AC P94226;
DT 01-MAY-1997 (TREMBLrel. 03, Created)
DT 01-MAY-1997 (TREMBLrel. 03, Last sequence update)
DT 01-JUN-2001 (TREMBLrel. 17, Last annotation update)
DE OUTER SURFACE PROTEIN C (FRAGMENT).
GN OSC.
OS Borrelia burgdorferi (Lyme disease spirochete).
OC Bacteria; Spirochaetales; Spirochaetaceae; Borrelia.
OX NCBI_TaxID=139;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=J1;
RX MEDLINE=96296448; PubMed=8709845;
RA Livey I., Gibbs C.P., Schuster R., Dornier F.;
RT "Evidence for lateral transfer and recombination in OspC variation in
Lyme disease Borrelia";
RL Mol. Microbiol. 18:257-269(1995).
DR EMBL; L42884; AAB36992.1;
DR InterPro: IPR001800; Lipoprotein_6.
DR Pfam: PF01441; Lipoprotein_6; 1.
DR ProDom: PD001149; Lipoprotein_6; 1.
FT NON_TER 1
FT 1
SQ SEQUENCE 193 AA; 20125 MW; DF3F926F8BF07290 CRC64;

Query Match 100.0%; Score 52; DB 2; Length 193;
Best Local Similarity 100.0%; Pred. No. 0.077;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 PVVAESPKKP 10
Db 184 PVVAESPKKP 193
|||||

RESULT 9
ID P94233 PRELIMINARY; PRT; 193 AA.
AC P94233;
DT 01-MAY-1997 (TREMBLrel. 03, Created)
DT 01-MAY-1997 (TREMBLrel. 03, Last sequence update)
DT 01-JUN-2001 (TREMBLrel. 17, Last annotation update)
DE OUTER SURFACE PROTEIN C (FRAGMENT).
GN OSC.
OS Borrelia burgdorferi (Lyme disease spirochete).
OC Bacteria; Spirochaetales; Spirochaetaceae; Borrelia.
OX NCBI_TaxID=139;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=ACAL1;
RX MEDLINE=96296448; PubMed=8709845;
RA Livey I., Gibbs C.P., Schuster R., Dornier F.;
RT "Evidence for lateral transfer and recombination in OspC variation in
Lyme disease Borrelia";
RL Mol. Microbiol. 18:257-269(1995).
DR EMBL; L42892; AAB37000.1;
DR InterPro: IPR001800; Lipoprotein_6.
DR Pfam: PF01441; Lipoprotein_6; 1.
DR ProDom: PD001149; Lipoprotein_6; 1.
FT NON_TER 1
FT 1
SQ SEQUENCE 193 AA; 20370 MW; CD52748E2EA2D36F CRC64;

Query Match 100.0%; Score 52; DB 2; Length 193;
Best Local Similarity 100.0%; Pred. No. 0.077;

Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 PVVAESPCKP 10
|||||
Db 184 PVVAESPCKP 193

RESULT 10

P94237 PRELIMINARY; PRT; 193 AA.
AC P94237;
DT 01-MAY-1997 (TREMblrel. 03, Created)
DT 01-MAY-1997 (TREMblrel. 03, Last sequence update)
DT 01-JUN-2001 (TREMblrel. 17, Last annotation update)
DE OUTER SURFACE PROTEIN C (FRAGMENT).
GN OSCP.
OS Borrelia burgdorferi (Lyme disease spirochete).
OC Bacteria; Spirochaetales; Spirochaetaceae; Borrelia.
OX NCBI_TaxID=139;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=J5015;
RX MEDLINE=96296448; PubMed=8709845;
RA Livey I., Gibbs C.P., Schuster R., Dorner F.;
RT "Evidence for lateral transfer and recombination in Oscp variation in
RL Lyme disease Borrelia.";
RL Mol. Microbiol. 18:257-269(1995).
DR InterPro: IPR001800; Lipoprotein_6.
DR Pfam: PF01441; Lipoprotein_6; 1.
DR Prodom: PD001149; Lipoprotein_6; 1.
FT NON_TER 1
FT SEQUENCE 193 AA; 20677 MW; C9500D959E13590D CRC64;

Query Match 100.0%; Score 52; DB 2; Length 193;
Best Local Similarity 100.0%; Pred. No. 0.077;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 PVVAESPCKP 10
|||||
Db 184 PVVAESPCKP 193

RESULT 11

P94247 PRELIMINARY; PRT; 194 AA.
AC P94247;
DT 01-MAY-1997 (TREMblrel. 03, Created)
DT 01-MAY-1997 (TREMblrel. 03, Last sequence update)
DT 01-JUN-2001 (TREMblrel. 17, Last annotation update)
DE OUTER SURFACE PROTEIN C (FRAGMENT).
GN OSCP.
OS Borrelia burgdorferi (Lyme disease spirochete).
OC Bacteria; Spirochaetales; Spirochaetaceae; Borrelia.
OX NCBI_TaxID=139;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=SIMON;
RX MEDLINE=96296448; PubMed=8709845;
RA Livey I., Gibbs C.P., Schuster R., Dorner F.;
RT "Evidence for lateral transfer and recombination in Oscp variation in
RL Lyme disease Borrelia.";
RL Mol. Microbiol. 18:257-269(1995).
DR InterPro: IPR001800; Lipoprotein_6.
DR Pfam: PF01441; Lipoprotein_6; 1.
DR Prodom: PD001149; Lipoprotein_6; 1.
FT NON_TER 1
FT SEQUENCE 194 AA; 20361 MW; EDC8E0F602E02DC9 CRC64;

Query Match 100.0%; Score 52; DB 2; Length 194;
Best Local Similarity 100.0%; Pred. No. 0.077;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 PVVAESPCKP 10
|||||
Db 185 PVVAESPCKP 194

RESULT 12

O9S3P4 PRELIMINARY; PRT; 194 AA.
AC O9S3P4;
DT 01-MAY-2000 (TREMblrel. 13, Created)
DT 01-MAY-2000 (TREMblrel. 13, Last sequence update)
DT 01-JUN-2001 (TREMblrel. 17, Last annotation update)
DE OUTER SURFACE PROTEIN C (FRAGMENT).
GN OSCP.
OS Borrelia burgdorferi (Lyme disease spirochete).
OC Bacteria; Spirochaetales; Spirochaetaceae; Borrelia.
OX NCBI_TaxID=139;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=J5B;
RX MEDLINE=96296448; PubMed=8709845;
RA Livey I., Gibbs C.P., Schuster R., Dorner F.;
RT "Evidence for lateral transfer and recombination in Oscp variation in
RL Lyme disease Borrelia.";
RL Mol. Microbiol. 18:257-269(1995).
DR InterPro: IPR001800; Lipoprotein_6.
DR Pfam: PF01441; Lipoprotein_6; 1.
DR Prodom: PD001149; Lipoprotein_6; 1.
FT NON_TER 1
FT SEQUENCE 194 AA; 20446 MW; CEEDC9FA5DF0D68F CRC64;

Query Match 100.0%; Score 52; DB 2; Length 194;
Best Local Similarity 100.0%; Pred. No. 0.077;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 PVVAESPCKP 10
|||||
Db 185 PVVAESPCKP 194

RESULT 13

P94229 PRELIMINARY; PRT; 194 AA.
AC P94229;
DT 01-MAY-1997 (TREMblrel. 03, Created)
DT 01-MAY-1997 (TREMblrel. 03, Last sequence update)
DT 01-JUN-2001 (TREMblrel. 17, Last annotation update)
DE OUTER SURFACE PROTEIN C (FRAGMENT).
GN OSCP.
OS Borrelia burgdorferi (Lyme disease spirochete).
OC Bacteria; Spirochaetales; Spirochaetaceae; Borrelia.
OX NCBI_TaxID=139;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=H9;
RX MEDLINE=96296448; PubMed=8709845;
RA Livey I., Gibbs C.P., Schuster R., Dorner F.;
RT "Evidence for lateral transfer and recombination in Oscp variation in
RL Lyme disease Borrelia.";
RL Mol. Microbiol. 18:257-269(1995).
DR InterPro: IPR001800; Lipoprotein_6.
DR Pfam: PF01441; Lipoprotein_6; 1.
DR Prodom: PD001149; Lipoprotein_6; 1.
FT NON_TER 1
FT SEQUENCE 194 AA; 20361 MW; EDC8E0F602E02DC9 CRC64;

FT NON_TER 194 194
SQ SEQUENCE 194 AA: 20270 MW: A016BB5336A9C981 CRC64;

Query Match
Best Local Similarity 100.0%; Score 52; DB 2; Length 194;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 PVVAESPCKP 10
Db 185 PVVAESPCKP 194

RESULT 14

O45030 PRELIMINARY; PRT; 194 AA.
AC O45030;
DT 01-NOV-1996 (TREMblrel. 01, Created)
DT 01-NOV-1996 (TREMblrel. 01, Last sequence update)
DT 01-DEC-2001 (TREMblrel. 19, Last annotation update)
DE OSPA PROTEIN (FRAGMENT).
OS OSPA.
GN Borrelia burgdorferi (Lyme disease spirochete).
OC Bacteria; Spirochaetales; Spirochaetaceae; Borrelia.
OX NCBI_Taxid=139;
RN (1)
RP SEQUENCE FROM N.A.
RC STRAIN=PLE;
RX MEDLINE=95395018; PubMed=7665660;
RA Jauris-Helpke S., Liegl G., Preac-Mursic V., Roessler D., Schwab E.,
RT Soutschek E., Will G., Wilske B.,
RT "Molecular analysis of genes encoding outer surface protein C (OspC)
of Borrelia burgdorferi sensu lato: relationship to ospA genotype and
evidence of lateral gene exchange of ospC".
RL J. Clin. Microbiol. 33:1860-1866(1995).
DR EMBL: X80255; CA56548.1; -
DR InterPro: IPR001800; Lipoprotein_6.
DR Pfam: PF01441; Lipoprotein_6; 1.
DR ProDom: PD001149; Lipoprotein_6; 1.
FT NON_TER 1 1
FT NON_TER 194 194
SQ SEQUENCE 194 AA: 20523 MW: 11D409DBBFD23288 CRC64;

Query Match
Best Local Similarity 100.0%; Score 52; DB 2; Length 194;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 PVVAESPCKP 10
Db 185 PVVAESPCKP 194

RESULT 15

O45187 PRELIMINARY; PRT; 207 AA.
AC O45187;
DT 01-NOV-1996 (TREMblrel. 01, Created)
DT 01-NOV-1996 (TREMblrel. 01, Last sequence update)
DT 01-DEC-2001 (TREMblrel. 18, Last annotation update)
DE OUTER SURFACE PROTEIN C PRECURSOR.
OS OSCPC.
GN Borrelia garinii.
OC Bacteria; Spirochaetales; Spirochaetaceae; Borrelia.
OX NCBI_Taxid=29519;
RN (1)
RP SEQUENCE FROM N.A.
RC STRAIN=PTROB;
RX MEDLINE=95213332; PubMed=7699024;
RA Wilske B., Jauris-Helpke S., Lobentanzer R., Pradel T.,
RT Preac-Mursic V., Roessler D., Soutschek E., Johnson R.C.,
RT "Phenotypic analysis of outer surface protein C (OspC) of Borrelia
burgdorferi sensu lato by monoclonal antibodies: relationship to

RT genospecies and OspA serotype.";
RL J. Clin. Microbiol. 33:103-109(1995).
RN (2)

RP SEQUENCE FROM N.A.

RC STRAIN=PTROB;
RX MEDLINE=95395018; PubMed=7665660;

RA Jauris-Helpke S., Liegl G., Preac-Mursic V., Roessler D., Schwab E.,
RT Soutschek E., Will G., Wilske B.,

RT "Molecular analysis of genes encoding outer surface protein C (OspC)
of Borrelia burgdorferi sensu lato: relationship to ospA genotype and
evidence of lateral gene exchange of ospC".
RL J. Clin. Microbiol. 33:1860-1866(1995).
RN (3)

RP SEQUENCE FROM N.A.
RC STRAIN=PCSF, PBAEII, PFII, PELK, PMUE, AND PSB;
RA Marconi R.T., Hohenberger S., Jauris-Helpke S., Schulte-Spechtel U.,
RA Lavole C.P., Roessler D., Wilske B.,

RT "Genetic analysis of B. garinii OspA-serotype 4 strains associated with
neuroborreliosis: evidence for extensive genetic homogeneity".
RL Submitted (FEB-1999) to the EMBL/Genbank/DBJ databases.
DR EMBL: X83554; CA58544.1; -
DR EMBL: AJ236908; CAB46238.1; -
DR EMBL: AJ132793; CAB46231.1; -
DR EMBL: AJ132796; CAB46234.1; -
DR EMBL: AJ132797; CAB46235.1; -
DR EMBL: AJ132798; CAB46236.1; -
DR EMBL: AJ236907; CAB46237.1; -
DR InterPro: IPR001800; Lipoprotein_6.
DR Pfam: PF01441; Lipoprotein_6; 1.
DR ProDom: PD001149; Lipoprotein_6; 1.
FT SIGNAL.
FT KW
FT SIGNAL

FT CHAIN 1 18 POTENTIAL.
FT SEQUENCE 19 207 POTENTIAL.
SQ SEQUENCE 207 AA: 22321 MW: 20889AA70E63A49D CRC64;

Query Match
Best Local Similarity 100.0%; Score 52; DB 2; Length 207;
Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 PVVAESPCKP 10
Db 198 PVVAESPCKP 207

Search completed: October 12, 2002, 20:47:11
Job time: 11965 sec